JC07 Rec'd PCT/PTO 24 APR 2001

FORM PTO-1390 (REV 5-93)	U.S DEPARTMENT OF COMMERCE PATENT AND TRADEMARK OFFICE	ATTORNEY DOCKET NO. PB370US
TRANSMITTAL LETTER TO THE UNITED STATES DESIGNATED/ELECTED OFFICE (DO/EO/US) CONCERNING A FILING UNDER 35 U.S.C. 371		DATE. April 24, 2001
		U S APPLN NO. (IF KNOWN, SEE 37 C F R 1.5) Unknown 09/830228

INTERNATIONAL APPLICATION NO PCT/US98/12764	INTERNATIONAL FILING DATE June 18, 1998	PRIORITY DATE CLAIMED June 20, 1997

TITLE OF INVENTION *Borrelia Burgdorferi* Polynucleotides and Sequences

APPLICANT(S) FOR DO/EO/US Human Genome Sciences, Inc et al.

1. ☒ This is a **FIRST** submission of items concerning a filing under 35 U S C 371. (THE BASIC FILING FEE IS ATTACHED)

2. ☐ This is a **SECOND** or **SUBSEQUENT** submission of items concerning a filing under 35 U S C 371

3. ☒ This express request to begin national examination procedures [35 U S C 371(f)] at any time rather than delay examination until the expiration of the applicable time limit set in 35 U.S C 371(b) and PCT Articles 22 and 39(1).

4. ☒ A proper demand for International Preliminary Amendment was made by the 19th month from the earliest claimed priority date.

5. ☒ A copy of the International Application as filed [35 U.S.C 371(c)(2)]
   a. ☒ is transmitted herewith (required only if not transmitted by the International Bureau)
   b. ☐ has been transmitted by the International Bureau
   c. ☐ is not required, as the application was filed in the United States Receiving Office (RO/US).

6. ☐ A translation of the International Application into English [35 U S C. 371(c)(2)]

7. ☐ Amendments to the claims of the International Application under PCT Article 19 [35 U.S C 371(c)(3)]
   a. ☐ are transmitted herewith (required only if not transmitted by the International Bureau)
   b. ☐ have been transmitted by the International Bureau
   c. ☐ have not been made, however, the time limit for making such amendments has NOT expired.
   d. ☐ have not been made and will not be made

8. ☐ A translation of the amendments to the claims under PCT Article 19 [35 U S.C 371(c)(3)].

9. ☒ An unexecuted oath or declaration of the inventor(s) [35 U S.C. 371(c)(4)]

10. ☐ A translation of the annexes to the International Preliminary Examination Report under PCT Article 36 [35 U S C 371(c)(5)].

Items 11 - 16 below concern other document(s) or information included:

11. ☒ An Information Disclosure Statement under 37 C F R. 1.97 and 1 98

12. ☐ An assignment document for recording. A separate cover sheet in compliance with 37 C.F R 3 28 and 3.31 is included.

13. ☐ A FIRST preliminary amendment.
    ☐ A SECOND or SUBSEQUENT preliminary amendment.

14. ☐ A substitute specification.

15. ☐ A change of power of attorney and/or address letter.

16. ☒ Other items or information ☒ Paper copy (on pages 156-1114) of Sequence Listing and Request under 37 CFR 1 821(e)
    CHECK NO. ☐
    Drawings (☐ sheets)

UNDER SEAL

JC18 Rec'd PCT/PTO 24 APR 2001

U.S. APPLN. NO. (IF KNOWN SEE 37 C.F.R. 1.5(b)) 09/830228	INTERNATIONAL APPLICATION NO. PCT/US98/12764	ATTORNEY DOCKET NO. PB370US DATE: April 24, 2001

17. ☒ The following fees are submitted.	CALCULATIONS	PTO USE ONLY
Basic National Fee [37 C.F.R. 1.492(a)(1)-(5)]:		
Search Report has been prepared by the EPO or JPO $860.00		
International preliminary examination fee paid to USPTO (37 C.F.R. 1.482) ... $690.00		
No international preliminary examination fee paid to USPTO (37 C.F.R. 1.482) but international search fee paid to USPTO [37 C.F.R. 1.445(a)(2)] ... $710.00		
Neither international preliminary examination fee (37 C.F.R. 1.482) or international search fee [37 C.F.R. 1.445(a)(2)] paid to USPTO ... $1,000.00		
International preliminary examination fee paid to USPTO (37 C.F.R. 1.482) and all claims satisfied provisions of PCT Article 33(2)-(4) $100.00		
ENTER APPROPRIATE BASIC FEE AMOUNT =	$690.00	
Surcharge of $130.00 for furnishing the oath or declaration later than ☐ 20 ☒ 30 months from the earliest claimed priority date [37 C.F.R. 1.492(e)]	$130.00	

Claims	Number Filed	Number Extra	Rate		
Total Claims	21 - 20 =	1	X $18.00	$18.00	
Independent Claims	6 - 3 =	3	X $80.00	$240.00	
Multiple dependent claim(s) (if applicable)			+ $270.00		
TOTAL OF ABOVE CALCULATIONS =				$1,078.00	
Reduction by one-half for filing by small entity, if applicable. Verified Small Entity statement must also be filed (Note 37 C.F.R. 1.9, 1.27, 1.28).			-	$	
SUBTOTAL =				$1,078.00	
Processing fee of $130.00 for furnishing the English translation later the ☐ 20 ☐ 30 months from the earliest claimed priority date [37 C.F.R. 1.492(f)].			+	$	
TOTAL NATIONAL FEE =				$1,078.00	
Fee for recording the enclosed assignment [37 C.F.R. 1.21(h)]. The assignment must be accompanied by an appropriate cover sheet (37 C.F.R. 3.28, 3.31) $40.00 per property			+	$	
TOTAL FEES ENCLOSED =				$1,078.00	
				Amount to be refunded	$
				Charged	$

a. ☐ A check in the amount of $ to cover the above fees is enclosed.
b. ☒ Please charge my Deposit Account No. 08-3425 in the amount of $1,582.00 to cover the above fee. A duplicate copy of this sheet is enclosed.
c. ☒ The Commissioner is hereby authorized to charge any additional fees which may be required, or credit any overpayment to Deposit Account No. 08-3425.

NOTE: Where an appropriate time limit under 37 C.F.R. 1.494 or 1.495 has not been met, a petition to revive [37 C.F.R. 1.137(a) or (b)] must be filed and granted to restore the application to pending status.

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IN THE UNITED STATES PATENT AND TRADEMARK OFFICE

Application of: Fraser et al.

Application Number: to be assigned

Group Art Unit: to be assigned

Filed: herewith

Examiner: to be assigned

Title: *Borrelia Burgdorferi* Polynucleotides and Sequences

Attny. Docket No. PB370US

REQUEST UNDER 37 C.F.R. § 1.821(e)

Assistant Commissioner for Patents
Washington, D.C. 20231

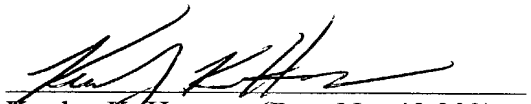
Sir :

The above-identified patent application is a US National Stage Entry application of International application Serial No. PCT/US98/12764, filed June 18, 1998. The sequence listing of the instant application is identical to the sequence content of the computer readable sequence listing filed in connection with application Serial No. PCT/US98/12764, on June 18, 1998.

In accordance with 37 C.F.R. § 1.821(e), please use the computer readable form filed in connection with application Serial No. PCT/US98/12764as the computer readable form for the instant application. It is understood that the Patent and Trademark Office will make the necessary change in application number and filing date for the computer readable form that will be used for the instant application. A paper copy of the Sequence Listing is included in the specification of the instant application filed herewith. Applicants hereby certify that the paper copy of the Sequence Listing filed herewith and the computer readable sequence listing filed on June 18, 1998 in connection with International application Serial No. PCT/US98/12764 are the same and do not include new matter.

Respectfully submitted

Date: April 24, 2001



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09/830228
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Borrelia burgdorferi Polynucleotides and Sequences

Field of the Invention

The present invention relates to the field of molecular biology. In particular, it relates to, among other things, nucleotide sequences of *Borrelia burgdorferi*, contigs, ORFs, fragments, probes, primers and related polynucleotides thereof, peptides and polypeptides encoded by the sequences, and uses of the polynucleotides and sequences thereof, such as in fermentation, polypeptide production, assays and pharmaceutical development, among others.

Statement as to Rights to Inventions Made Under Federally-Sponsored Research and Development

Part of the work performed during development of this invention utilized U.S. Government funds. The U.S. Government may have certain rights in the invention - DE-FC02-95ER61962; DE-FCO2-95ER61963; and NAGW 2554.

Background of the Invention

Spirochetes are a family of motile, unicellular, spiral-shaped bacteria which share a number of structural characteristics. Three genera of the spirochetes are pathogenic in humans: (a) *Treponema*, which includes the pathogens that cause syphilis (*T. pallidum*), yaws (*T. pertenue*), and pinta (*T. carateum*); (b) *Borrelia*, which includes the pathogens that cause epidemic and endemic relapsing fever and Lyme disease; and (c) *Leptospira*, which includes a wide variety of small spirochetes that cause mild to serious systemic human illness (Koff, A. B. and Rosen, T. *J. Am. Acad. Dermatol.* **29**:519-535 (1993)).

Lyme borreliosis, more commonly known as Lyme disease, is presently the most common human disease in the United States transmitted by an arthropod vector. Centers for Disease Control, Morbid. Mortal. Weekly Rep. 44:590-591 (1995). Further, infection of household pets, such as dogs, is a considerable problem. The causative agent of this affliction is the spirochete *Borrelia burgdorferi*, which is generally transmitted to mammalian hosts by feeding ticks. Barbour, A. and Fish, D. Science 260:1610-1616 (1993). Once the bacteria pass through the skin they disseminate and produce a variety of clinical manifestations. Diagnosis of this disease is often made serologically by the identification of antiborrelial antibodies. Hilton, E. et al., J. Clin. Microbiol. 35:774-776 (1997).

While initial symptoms often include a rash at the infection point, Lyme disease is a multisystemic disorder that may include arthritic, carditic, and neurological manifestations. While antibiotics are currently used to treat active cases of Lyme disease, *B. burgdorferi* appears to be able to persist even after prolonged antibiotic treatment. Further, *B. burgdorferi* can persist for years in a mammalian host even in the presence of an active immune response. Straubinger, R. et al., J. Clin. Microbiol. 35:111-116 (1997); Steere, A., N. Engl. J. Med. 321:586-596 (1989).

Animal models have proven useful for studying the progression of Lyme disease, methods for preventing this disease, and immunological responses to antigenic challenges with *B. burgdorferi* proteins. Garcia-Monoco, J. et al., J. Infect. Dis. 175:1243-1245 (1997). Using a canine model, Starubinger, R. et al., Infect. Immun. 65:1273-1285 (1977), demonstrated that *B. burgdorferi* migrates into joints and induces up-regulation of interleukin-8 in synovial membranes. Similarly, *B. burgdorferi* induction of interleukin-8 production has been demonstrated in cultured human endothelial cells. Burns, M. et al., Infect. Immun. 65:1217-1222 (1997).

Antigenic heterogeneity has been postulated as a mechanism used by *B. burgdorferi* for evasion of host immune responses. Schwan, T. et al., Can. J. Microbiol. 37:450-454 (1991). In support of this mechanism, antigenic variation has been described with other pathogenic bacteria. Hagbloom, P. et al., Nature 315:156-158 (1985). Further, cassette type genetic recombination of genes encoding *B. burgdorferi* surface proteins has been shown to decrease the antigenicity of these organisms to antibodies generated against strains which have not undone the same recombination. Zhang, J. et al., Cell 89:275-285 (1997).

A number of different types of Lyme disease vaccines have been tested and shown to induce immunological responses. Whole-cell *B. burgdorferi* vaccines have been shown to induce both immunological responses and protective immunity in several animal models. Reviewed in Wormser, G., Clin. Infect. Dis. 21:1267-1274 (1995). For example, dogs inoculated with a chemically inactivated whole-cell vaccine primarily develop antibodies to outer surface membrane proteins of the administered organism. Further, passive immunity has been also demonstrated in animals using *B. burgdorferi* specific antisera. Similarly, passive immunity is conferred human by the administration of sera obtained from Lyme disease patients.

While whole-cell Lyme disease vaccines confer protective immunity in animal models, use of such vaccines presents the risk that responsive antibodies will be generated which cross react with human antigens. Reviewed in Wormser, G., supra. This problem is at least partly the result of the production of *B. burgdorferi* specific antibodies which cross-react with hepatocytes and both muscle and nerve cells. *B. burgdorferi* heat shock proteins and the 41-kd flagellin subunit are believed to contain the antigens against which these cross-reactive antibodies are generated.

It is clear that the etiology of diseases mediated or exacerbated by *B. burgdorferi* genes, and that characterizing the genes and their patterns of expression would add dramatically to our

understanding of the organism and its host interactions. Knowledge of *B. burgdorferi* genes and genomic organization would dramatically improve understanding of disease etiology and lead to improved and new ways of preventing, ameliorating, arresting and reversing diseases. Moreover, characterized genes and genomic fragments of *B. burgdorferi* would provide reagents for, among other things, detecting, characterizing and controlling *B. burgdorferi* infections. There is a need therefore to characterize the genome of *B. burgdorferi* and for polynucleotides and sequences of this organism.

SUMMARY OF THE INVENTION

The present invention is based on the sequencing of fragments of the *Borrelia burgdorferi* genome. The primary nucleotide sequences which were generated are provided in SEQ ID NOS:1-155.

The present invention provides the complete nucleotide sequence of the *Borrelia burgdorferi* chromosome and 154 contigs representing the majority of the sequence of the B. burgdorferi extrachromosomal elements, all of which are listed in tables below and set out in the Sequence Listing submitted herewith, and representative fragments thereof, in a form which can be readily used, analyzed, and interpreted by a skilled artisan. In one embodiment, the present invention is provided as contiguous strings of primary sequence information corresponding to the nucleotide sequences depicted in SEQ ID NOS: 1-155.

The present invention further provides nucleotide sequences which are at least 95%, 96%, 97%, 98%, and 99%, identical to the nucleotide sequences of SEQ ID NOS:1-155, ORF IDs and corresponding ORFs.

The nucleotide sequences of SEQ ID NOS:1-155, ORF ID or ORF within, a representative fragment thereof, or a nucleotide sequence which is at least 95% identical to said nucleotide sequence may be provided in a variety of mediums to facilitate its use. In one application of this embodiment, the sequences of the present invention are recorded on computer readable media. Such media includes, but is not limited to: magnetic storage media, such as floppy discs, hard disc storage medium, and magnetic tape; optical storage media such as CD-ROM; electrical storage media such as RAM and ROM; and hybrids of these categories such as magnetic/optical storage media.

The present invention further provides systems, particularly computer-based systems which contain the sequence information herein described stored in a data storage means. Such systems are designed to identify commercially important fragments of the *Borrelia burgdorferi* genome.

Another embodiment of the present invention is directed to fragments of the *Borrelia burgdorferi* genome having particular structural or functional attributes. Such fragments of the *Borrelia burgdorferi* genome of the present invention include, but are not limited to, fragments which encode peptides, hereinafter referred to as open reading frames or ORFs, fragments which modulate the expression of an operably linked ORF, hereinafter referred to as expression

modulating fragments or EMFs, and fragments which can be used to diagnose the presence of *Borrelia burgdorferi* in a sample, hereinafter referred to as diagnostic fragments or DFs.

Each of the ORF IDs and ORFs in fragments of the *Borrelia burgdorferi* genome disclosed in Tables 1-6, and the EMFs found 5' prime of the initiation codon, can be used in numerous ways as polynucleotide reagents. For instance, the sequences can be used as diagnostic probes or amplification primers for detecting or determining the presence of a specific microbe in a sample, to selectively control gene expression in a host and in the production of polypeptides, such as polypeptides encoded by ORFs of the present invention, particular those polypeptides that have a pharmacological activity.

The present invention further includes recombinant constructs comprising one or more fragments of the *Borrelia burgdorferi* genome of the present invention. The recombinant constructs of the present invention comprise vectors, such as a plasmid or viral vector, into which a fragment of the *Borrelia burgdorferi* has been inserted.

The present invention further provides host cells containing any of the isolated fragments of the *Borrelia burgdorferi* genome of the present invention. The host cells can be a higher eukaryotic host cell, such as a mammalian cell, a lower eukaryotic cell, such as a yeast cell, or a procaryotic cell such as a bacterial cell.

The present invention is further directed to isolated polypeptides and proteins encoded by ORFs of the present invention. A variety of methods, well known to those of skill in the art, routinely may be utilized to obtain any of the polypeptides and proteins of the present invention. For instance, polypeptides and proteins of the present invention having relatively short, simple amino acid sequences readily can be synthesized using commercially available automated peptide synthesizers. Polypeptides and proteins of the present invention also may be purified from bacterial cells which naturally produce the protein. Yet another alternative is to purify polypeptide and proteins of the present invention from cells which have been altered to express them.

The invention further provides methods of obtaining homologs of the fragments of the *Borrelia burgdorferi* genome of the present invention and homologs of the proteins encoded by the ORFs of the present invention. Specifically, by using the nucleotide and amino acid sequences disclosed herein as a probe or as primers, and techniques such as PCR cloning and colony/plaque hybridization, one skilled in the art can obtain homologs.

The invention further provides antibodies which selectively bind polypeptides and proteins of the present invention. Such antibodies include both monoclonal and polyclonal antibodies.

The invention further provides hybridomas which produce the above-described antibodies. A hybridoma is an immortalized cell line which is capable of secreting a specific monoclonal antibody.

The present invention further provides methods of identifying test samples derived from cells which express one of the ORFs of the present invention, or a homolog thereof. Such

methods comprise incubating a test sample with one or more of the antibodies of the present invention, or one or more of the DFs of the present invention, under conditions which allow a skilled artisan to determine if the sample contains the ORF or product produced therefrom.

In another embodiment of the present invention, kits are provided which contain the necessary reagents to carry out the above-described assays.

Specifically, the invention provides a compartmentalized kit to receive, in close confinement, one or more containers which comprises: (a) a first container comprising one of the antibodies, or one of the DFs of the present invention; and (b) one or more other containers comprising one or more of the following: wash reagents, reagents capable of detecting presence of bound antibodies or hybridized DFs.

Using the isolated proteins of the present invention, the present invention further provides methods of obtaining and identifying agents capable of binding to a polypeptide or protein encoded by one of the ORFs of the present invention. Specifically, such agents include, as further described below, antibodies, peptides, carbohydrates, pharmaceutical agents and the like. Such methods comprise steps of: (a)contacting an agent with an isolated protein encoded by one of the ORFs of the present invention; and (b)determining whether the agent binds to said protein.

The present genomic sequences of *Borrelia burgdorferi* will be of great value to all laboratories working with this organism and for a variety of commercial purposes. Many fragments of the *Borrelia burgdorferi* genome will be immediately identified by similarity searches against GenBank or protein databases and will be of immediate value to *Borrelia burgdorferi* researchers and for immediate commercial value for the production of proteins or to control gene expression.

The methodology and technology for elucidating extensive genomic sequences of bacterial and other genomes has and will greatly enhance the ability to analyze and understand chromosomal organization. In particular, sequenced contigs and genomes will provide the models for developing tools for the analysis of chromosome structure and function, including the ability to identify genes within large segments of genomic DNA, the structure, position, and spacing of regulatory elements, the identification of genes with potential industrial applications, and the ability to do comparative genomic and molecular phylogeny.

DESCRIPTION OF THE FIGURES

FIGURE 1 is a block diagram of a computer system (102) that can be used to implement computer-based systems of present invention.

FIGURE 2 is a schematic diagram depicting the data flow and computer programs used to collect, assemble, edit and annotate the contigs of the *Borrelia burgdorferi* genome of the present invention. Both Macintosh and Unix platforms are used to handle the AB 373 and 377 sequence data files, largely as described in Kerlavage *et al.*, *Proceedings of the Twenty-Sixth*

Annual Hawaii International Conference on System Sciences, 585, IEEE Computer Society Press, Washington D.C. (1993). Factura (AB) is a Macintosh program designed for automatic vector sequence removal and end-trimming of sequence files. The program Loadis runs on a Macintosh platform and parses the feature data extracted from the sequence files by Factura to the Unix based *Borrelia burgdorferi* relational database. Assembly of contigs (and whole genome sequences) is accomplished by retrieving a specific set of sequence files and their associated features using Extrseq, a Unix utility for retrieving sequences from an SQL database. The resulting sequence file is processed to trim portions of the sequences with a high rate ambiguous nucleotides. The sequence files were assembled using TIGR Assembler, an assembly engine designed at The Institute for Genomic Research (TIGR) for rapid and accurate assembly of thousands of sequence fragments. The collection of contigs generated by the assembly step is loaded into the database with the lassie program. Identification of open reading frames (ORFs) is accomplished by processing contigs with zorf. The ORFs are searched against *B. burgdorferi* sequences from GenBank and against all protein sequences using the BLASTN and BLASTP programs, described in Altschul *et al.*, *J. Mol. Biol. 215:* 403-410 (1990). Results of the ORF determination and similarity searching steps were loaded into the database. As described below, some results of the determination and the searches are set out in Tables 1-6.

DETAILED DESCRIPTION OF ILLUSTRATIVE EMBODIMENTS

The present invention is based on the sequencing of fragments of the *Borrelia burgdorferi* genome and analysis of the sequences. The primary nucleotide sequences generated by sequencing the fragments are provided in SEQ ID NOS: 1-155. (As used herein, the "primary sequence" refers to the nucleotide sequence represented by the IUPAC nomenclature system.) SEQ ID NOS:1-155

In addition, the present invention provides the nucleotide sequences of SEQ ID NOS: 1-155, or representative fragments thereof, in a form which can be readily used, analyzed, and interpreted by a skilled artisan.

As used herein, a "representative fragment of the nucleotide sequence depicted in SEQ ID NOS:1-155" refers to any portion of the SEQ ID NOS: 1-155 which is not presently represented within a publicly available database. Preferred representative fragments of the present invention are *Borrelia burgdorferi* open reading frames (ORFs) represented by ORF IDs, expression modulating fragments (EMFs) and diagnostic fragments (DFs)which can be used to diagnose the presence of *Borrelia burgdorferi* in sample. A non-limiting identification of preferred representative portions are provided in Tables 1-6 as ORF IDs. As discussed in detail below, the information provided in SEQ ID NOS:1-155 and in Tables 1-6 together with routine cloning, synthesis, sequencing and assay methods will enable those skilled in the art to clone and sequence all "representative fragments" of interest, including ORFs encoding a large variety of *Borrelia burgdorferi* proteins.

The present invention is further directed to nucleic acid molecules encoding portions or fragments of the nucleotide sequences described herein. Fragments include portions of the nucleotide sequences of Table 1-6 (ORF IDs) and SEQ ID NOS:1-155, at least 10 contiguous nucleotides in length selected from any two integers, one of which representing a 5' nucleotide position and a second of which representing a 3' nucleotide position, where the first nucleotide for each nucleotide sequence in SEQ ID NOS:1-155 is position 1 (therefore, the sequence postions for each ORF ID is determined by the numbering of the SEQ ID comprising the ORF ID). That is, every combination of a 5' and 3' nucleotide position that a fragment at least 10 contiguous nucleotides in length could occupy is included in the invention. At least means a fragment may be 10 contiguous nucleotide bases in length or any integer between 10 and the length of an entire nucleotide sequence of SEQ ID NOS:1-155 minus 1. Therefore, included in the invention are contiguous fragments specified by any 5' and 3' nucleotide base positions of a nucleotide sequences of SEQ ID NOS:1-155 wherein the contiguous fragment is any integer between 10 and the length of an entire nucleotide sequence minus 1.

Further, the invention includes polynucleotides comprising fragments specified by size, in nucleotides, rather than by nucleotide positions. The invention includes any fragment size, in contiguous nucleotides, selected from integers between 10 and the length of an entire ORF ID or SEQ ID NO:, minus 1. Preferred sizes of contiguous nucleotide fragments include 20 nucleotides, 30 nucleotides, 40 nucleotides, 50 nucleotides. Other preferred sizes of contiguous nucleotide fragments, which may be useful as diagnostic probes and primers, include fragments 50-300 nucleotides in length which include, as discussed above, fragment sizes representing each integer between 50-300. Larger fragments are also useful according to the present invention corresponding to most, if not all, of the nucleotide sequences shown in Tables 1-6 (ORF IDs) and SEQ ID NOS:1-155. The preferred sizes are, of course, meant to exemplify not limit the present invention as all size fragments, representing any integer between 10 and the length of an entire nucleotide sequence minus 1, of each ORF ID and SEQ ID NO:, are included in the invention.

The present invention also provides for the exclusion of any fragment, specified by 5' and 3' base positions or by size in nucleotide bases as described above for any ORF ID or SEQ ID NOS:1-155. Any number of fragments of nucleotide sequences in ORF IDs or SEQ ID NOS:1-155, specified by 5' and 3' base positions or by size in nucleotides, as described above, may be excluded from the present invention.

While the presently disclosed sequences of SEQ ID NOS: 1-155 are highly accurate, sequencing techniques are not perfect and, in relatively rare instances, further investigation of a fragment or sequence of the invention may reveal a nucleotide sequence error present in a nucleotide sequence disclosed in SEQ ID NOS: 1-155. However, once the present invention is made available (*i.e.*, once the information in SEQ ID NOS: 1-155 and Tables 1-6 has been made available), resolving a rare sequencing error in SEQ ID NOS: 1-155 will be well within the skill

of the art. The present disclosure makes available sufficient sequence information to allow any of the described contigs or portions thereof to be obtained readily by straightforward application of routine techniques. Further sequencing of such polynucleotide may proceed in like manner using manual and automated sequencing methods which are employed ubiquitous in the art. Nucleotide sequence editing software is publicly available. For example, Applied Biosystem's (AB) AutoAssembler can be used as an aid during visual inspection of nucleotide sequences. By employing such routine techniques potential errors readily may be identified and the correct sequence then may be ascertained by targeting further sequencing effort, also of a routine nature, to the region containing the potential error.

Even if all of the very rare sequencing errors in SEQ ID NOS: 1-155 were corrected, the resulting nucleotide sequences would still be at least 95% identical, nearly all would be at least 99% identical, and the great majority would be at least 99.9% identical to the nucleotide sequences of SEQ ID NOS: 1-155.

As discussed elsewhere herein, polynucleotides of the present invention readily may be obtained by routine application of well known and standard procedures for cloning and sequencing DNA. Detailed methods for obtaining libraries and for sequencing are provided below, for instance. A wide variety of *Borrelia burgdorferi* strains that can be used to prepare *B. burgdorferi* genomic DNA for cloning and for obtaining polynucleotides of the present invention are available to the public from recognized depository institutions, such as the American Type Culture Collection (ATCC). While the present invention is enabled by the sequences and other information herein disclosed, the *B. burgdorferi* strain that provided the DNA of the present Sequence Listing, has been deposited with the ATCC, 10801 University Blvd. Manassas, VA 20110-2209, as Deposit No. 202012, on 8 August 1997. The ATCC Deposit is provided merely as a convenience to those of skill in the art. Reference to the deposit is not a waiver of any rights of the inventors or their assignees in the present subject matter.

The nucleotide sequences of the genomes from different strains of *Borrelia burgdorferi* differ somewhat. However, the nucleotide sequences of the genomes of all *Borrelia burgdorferi* strains will be at least 95% identical, in corresponding part, to the nucleotide sequences provided in SEQ ID NOS: 1-155 and the ORF IDs within. Nearly all will be at least 99% identical and the great majority will be 99.9% identical.

The present application is further directed to nucleic acid molecules at least 90%, 95%, 96%, 97%, 98% or 99% identical to a nucleic acid sequence shown in SEQ ID NOS: 1-155 and the ORF IDs within. The above nucleic acid sequences are included irrespective of whether they encode a polypeptide having *B. burgdorferi* activity. This is because even where a particular nucleic acid molecule does not encode a polypeptide having *B. burgdorferi* activity, one of skill in the art would still know how to use the nucleic acid molecule, for instance, as a hybridization probe. Uses of the nucleic acid molecules of the present invention that do not encode a polypeptide having *B. burgdorferi* activity include, *inter alia*, isolating a *B. burgdorferi* gene or allelic variants thereof from a DNA library, and detecting *B. burgdorferi* mRNA expression from

biological or environmental samples, suspected of containing *B. burgdorferi* by Northern Blot, PCR, or similar analysis.

Preferred, are nucleic acid molecules having sequences at least 90%, 95%, 96%, 97%, 98% or 99% identical to the nucleic acid sequence shown in SEQ ID NOS: 1-155, the ORF IDs, and the ORF within each ORF ID, which do, in fact, encode a polypeptide having *B. burgdorferi* protein activity. By "a polypeptide having *B. burgdorferi* activity" is intended polypeptides exhibiting activity similar, but not necessarily identical, to an activity of the *B. burgdorferi* protein of the invention, as measured in a particular biological assay suitable for measuring activity of the specified protein.

Due to the degeneracy of the genetic code, one of ordinary skill in the art will immediately recognize that a large number of the nucleic acid molecules having a sequence at least 90%, 95%, 96%, 97%, 98%, or 99% identical to the nucleic acid sequences shown in SEQ ID NOS: 1-155, the ORF IDs, and the ORF within each ORF ID, will encode a polypeptide having *B. burgdorferi* protein activity. In fact, since degenerate variants of these nucleotide sequences all encode the same polypeptide, this will be clear to the skilled artisan even without performing the above described comparison assay. It will be further recognized in the art that, for such nucleic acid molecules that are not degenerate variants, a reasonable number will also encode a polypeptide having *B. burgdorferi* protein activity. This is because the skilled artisan is fully aware of amino acid substitutions that are either less likely or not likely to significantly effect protein function (e.g., replacing one aliphatic amino acid with a second aliphatic amino acid), as further described below.

The biological activity or function of the polypeptides of the present invention are expected to be similar or identical to polypeptides from other bacteria that share a high degree of structural identity/similarity. Tables 1, 2, 4, and 5 lists accession numbers and descriptions for the closest matching sequences of polypeptides available through Genbank. It is therefore expected that the biological activity or function of the polypeptides of the present invention will be similar or identical to those polypeptides from other bacterial genuses, species, or strains listed in Tables 1, 2, 4, and 5.

By a polynucleotide having a nucleotide sequence at least, for example, 95% "identical" to a reference nucleotide sequence of the present invention, it is intended that the nucleotide sequence of the polynucleotide is identical to the reference sequence except that the polynucleotide sequence may include up to five point mutations per each 100 nucleotides of the reference nucleotide sequence encoding the *B. burgdorferi* polypeptide. In other words, to obtain a polynucleotide having a nucleotide sequence at least 95% identical to a reference nucleotide sequence, up to 5% of the nucleotides in the reference sequence may be deleted, inserted, or substituted with another nucleotide. The query sequence may be an entire sequence shown in SEQ ID NOS: 1-155, an ORF ID, or the ORF within each ORF ID, or any fragment specified as described herein.

As a practical matter, whether any particular nucleic acid molecule or polypeptide is at least 90%, 95%, 96%, 97%, 98% or 99% identical to a nucleotide sequence of the presence invention can be determined conventionally using known computer programs. A preferred method for determining the best overall match between a query sequence (a sequence of the present invention) and a subject sequence, also referred to as a global sequence alignment, can be determined using the FASTDB computer program based on the algorithm of Brutlag et al. *See* Brutlag et al. (1990) Comp. App. Biosci. 6:237-245. In a sequence alignment the query and subject sequences are both DNA sequences. An RNA sequence can be compared by first converting U's to T's. The result of said global sequence alignment is in percent identity. Preferred parameters used in a FASTDB alignment of DNA sequences to calculate percent identity are: Matrix=Unitary, k-tuple=4, Mismatch Penalty=1, Joining Penalty=30, Randomization Group Length=0, Cutoff Score=1, Gap Penalty=5, Gap Size Penalty=0.05, Window Size=500 or the lenght of the subject nucleotide sequence, whichever is shorter.

If the subject sequence is shorter than the query sequence because of 5' or 3' deletions, not because of internal deletions, a manual correction must be made to the results. This is because the FASTDB program does not account for 5' and 3' truncations of the subject sequence when calculating percent identity. For subject sequences truncated at the 5' or 3' ends, relative to the query sequence, the percent identity is corrected by calculating the number of bases of the query sequence that are 5' and 3' of the subject sequence, which are not matched/aligned, as a percent of the total bases of the query sequence. Whether a nucleotide is matched/aligned is determined by results of the FASTDB sequence alignment. This percentage is then subtracted from the percent identity, calculated by the above FASTDB program using the specified parameters, to arrive at a final percent identity score. This corrected score is what is used for the purposes of the present invention. Only nucleotides outside the 5' and 3' nucleotides of the subject sequence, as displayed by the FASTDB alignment, which are not matched/aligned with the query sequence, are calculated for the purposes of manually adjusting the percent identity score.

For example, a 90 nucleotide subject sequence is aligned to a 100 nucleotide query sequence to determine percent identity. The deletions occur at the 5' end of the subject sequence and therefore, the FASTDB alignment does not show a matched/alignment of the first 10 nucleotides at 5' end. The 10 unpaired nucleotides represent 10% of the sequence (number of nucleotides at the 5' and 3' ends not matched/total number of nucleotides in the query sequence) so 10% is subtracted from the percent identity score calculated by the FASTDB program. If the remaining 90 nucleotides were perfectly matched the final percent identity would be 90%. In another example, a 90 nucleotide subject sequence is compared with a 100 nucleotide query sequence. This time the deletions are internal deletions so that there are no nucleotides on the 5' or 3' of the subject sequence which are not matched/aligned with the query. In this case the percent identity calculated by FASTDB is not manually corrected. Once again, only nucleotides 5' and 3' of the

subject sequence which are not matched/aligned with the query sequence are manually corrected for. No other manual corrections are to made for the purposes of the present invention.

COMPUTER RELATED EMBODIMENTS

The nucleotide sequences provided in SEQ ID NOS: 1-155, including ORF IDs and corresponding ORFs, a representative fragment thereof, or a nucleotide sequence at least 95%, preferably at least 96%, 97%, 98% or 99%, and most preferably at least 99.9% identical to said nucleotide sequences may be "provided" in a variety of mediums to facilitate use thereof. As used herein, provided refers to a manufacture, other than an isolated nucleic acid molecule, which contains a nucleotide sequence of the present invention, a representative fragment thereof, or a nucleotide sequence at least 95%, preferably at least 99% and most preferably at least 99.9% identical to a polynucleotide of the present invention. Such a manufacture provides a large portion of the *Borrelia burgdorferi* genome and parts thereof (*e.g.*, a *Borrelia burgdorferi* open reading frame (ORF)) in a form which allows a skilled artisan to examine the manufacture using means not directly applicable to examining the *Borrelia burgdorferi* genome or a subset thereof as it exists in nature or in purified form.

In one application of this embodiment, a nucleotide sequence of the present invention can be recorded on computer readable media. As used herein, "computer readable media" refers to any medium which can be read and accessed directly by a computer. Such media include, but are not limited to: magnetic storage media, such as floppy discs, hard disc storage medium, and magnetic tape; optical storage media such as CD- ROM; electrical storage media such as RAM and ROM; and hybrids of these categories, such as magnetic/optical storage media. A skilled artisan can readily appreciate how any of the presently known computer readable mediums can be used to create a manufacture comprising computer readable medium having recorded thereon a nucleotide sequence of the present invention. Likewise, it will be clear to those of skill how additional computer readable media that may be developed also can be used to create analogous manufactures having recorded thereon a nucleotide sequence of the present invention.

As used herein, "recorded" refers to a process for storing information on computer readable medium. A skilled artisan can readily adopt any of the presently know methods for recording information on computer readable medium to generate manufactures comprising the nucleotide sequence information of the present invention.

A variety of data storage structures are available to a skilled artisan for creating a computer readable medium having recorded thereon a nucleotide sequence of the present invention. The choice of the data storage structure will generally be based on the means chosen to access the stored information. In addition, a variety of data processor programs and formats can be used to store the nucleotide sequence information of the present invention on computer readable medium. The sequence information can be represented in a word processing text file, formatted in commercially- available software such as WordPerfect and MicroSoft Word, or represented in the form of an ASCII file, stored in a database application, such as DB2, Sybase,

Oracle, or the like. A skilled artisan can readily adapt any number of data-processor structuring formats (*e.g.*, text file or database) in order to obtain computer readable medium having recorded thereon the nucleotide sequence information of the present invention.

Computer software is publicly available which allows a skilled artisan to access sequence information provided in a computer readable medium. Thus, by providing in computer readable form the nucleotide sequences of the present invention (e.g. SEQ ID NOS: 1-155), a representative fragment thereof, or a nucleotide sequence at least 95%, preferably at least 96%, 97%, 98%, 99% and most preferably at least 99.9% identical to a sequence of the present invention (e.g. SEQ ID NOS: 1-155) enables the skilled artisan routinely to access the provided sequence information for a wide variety of purposes.

The examples which follow demonstrate how software which implements the BLAST (Altschul *et al.*, *J. Mol. Biol. 215*:403-410 (1990)) and BLAZE (Brutlag *et al.*, *Comp. Chem. 17*:203-207 (1993)) search algorithms on a Sybase system was used to identify open reading frames (ORFs) within the *Borrelia burgdorferi* genome which contain homology to ORFs or proteins from both *Borrelia burgdorferi* and from other organisms. Among the ORFs discussed herein are protein encoding fragments of the *Borrelia burgdorferi* genome useful in producing commercially important proteins, such as enzymes used in fermentation reactions and in the production of commercially useful metabolites.

The present invention further provides systems, particularly computer-based systems, which contain the sequence information described herein. Such systems are designed to identify, among other things, commercially important fragments of the *Borrelia burgdorferi* genome.

As used herein, "a computer-based system" refers to the hardware means, software means, and data storage means used to analyze the nucleotide sequence information of the present invention. The minimum hardware means of the computer-based systems of the present invention comprises a central processing unit (CPU), input means, output means, and data storage means. A skilled artisan can readily appreciate that any one of the currently available computer-based system are suitable for use in the present invention.

As stated above, the computer-based systems of the present invention comprise a data storage means having stored therein a nucleotide sequence of the present invention and the necessary hardware means and software means for supporting and implementing a search means.

As used herein, "data storage means" refers to memory which can store nucleotide sequence information of the present invention, or a memory access means which can access manufactures having recorded thereon the nucleotide sequence information of the present invention.

As used herein, "search means" refers to one or more programs which are implemented on the computer-based system to compare a target sequence or target structural motif with the sequence information stored within the data storage means. Search means are used to identify fragments or regions of the present genomic sequences which match a particular target sequence or target motif. A variety of known algorithms are disclosed publicly and a variety of

commercially available software for conducting search means are and can be used in the computer-based systems of the present invention. Examples of such software includes, but is not limited to, MacPattern (EMBL), BLASTN and BLASTX (NCBIA). A skilled artisan can readily recognize that any one of the available algorithms or implementing software packages for conducting homology searches can be adapted for use in the present computer-based systems.

As used herein, a "target sequence" can be any DNA or amino acid sequence of six or more nucleotides or two or more amino acids. A skilled artisan can readily recognize that the longer a target sequence is, the less likely a target sequence will be present as a random occurrence in the database. The most preferred sequence length of a target sequence is from about 10 to 100 amino acids or from about 30 to 300 nucleotide residues. However, it is well recognized that searches for commercially important fragments, such as sequence fragments involved in gene expression and protein processing, may be of shorter length.

As used herein, "a target structural motif," or "target motif," refers to any rationally selected sequence or combination of sequences in which the sequence(s) are chosen based on a three-dimensional configuration which is formed upon the folding of the target motif. There are a variety of target motifs known in the art. Protein target motifs include, but are not limited to, enzymic active sites and signal sequences. Nucleic acid target motifs include, but are not limited to, promoter sequences, hairpin structures and inducible expression elements (protein binding sequences).

A variety of structural formats for the input and output means can be used to input and output the information in the computer-based systems of the present invention. A preferred format for an output means ranks fragments of the *Borrelia burgdorferi* genomic sequences possessing varying degrees of homology to the target sequence or target motif. Such presentation provides a skilled artisan with a ranking of sequences which contain various amounts of the target sequence or target motif and identifies the degree of homology contained in the identified fragment.

A variety of comparing means can be used to compare a target sequence or target motif with the data storage means to identify sequence fragments of the *Borrelia burgdorferi* genome. In the present examples, implementing software which implement the BLAST and BLAZE algorithms, described in Altschul *et al.*, *J. Mol. Biol. 215:* 403-410 (1990), is used to identify open reading frames within the *Borrelia burgdorferi* genome. A skilled artisan can readily recognize that any one of the publicly available homology search programs can be used as the search means for the computer-based systems of the present invention. Of course, suitable proprietary systems that may be known to those of skill also may be employed in this regard.

Figure 1 provides a block diagram of a computer system illustrative of embodiments of this aspect of present invention. The computer system 102 includes a processor 106 connected to a bus 104. Also connected to the bus 104 are a main memory 108 (preferably implemented as random access memory, RAM) and a variety of secondary storage devices 110, such as a hard drive 112 and a removable medium storage device 114. The removable medium storage device

114 may represent, for example, a floppy disk drive, a CD-ROM drive, a magnetic tape drive, *etc.* A removable storage medium 116 (such as a floppy disk, a compact disk, a magnetic tape, *etc.*) containing control logic and/or data recorded therein may be inserted into the removable medium storage device 114. The computer system 102 includes appropriate software for reading the control logic and/or the data from the removable medium storage device 114, once it is inserted into the removable medium storage device 114.

A nucleotide sequence of the present invention may be stored in a well known manner in the main memory 108, any of the secondary storage devices 110, and/or a removable storage medium 116. During execution, software for accessing and processing the genomic sequence (such as search tools, comparing tools, *etc.*) reside in main memory 108, in accordance with the requirements and operating parameters of the operating system, the hardware system and the software program or programs.

BIOCHEMICAL EMBODIMENTS

Other embodiments of the present invention are directed to isolated fragments of the *Borrelia burgdorferi* genome. The fragments of the *Borrelia burgdorferi* genome of the present invention include, but are not limited to fragments which encode peptides, hereinafter open reading frames (ORFs), fragments which modulate the expression of an operably linked ORF, hereinafter expression modulating fragments (EMFs) and fragments which can be used to diagnose the presence of *Borrelia burgdorferi* in a sample, hereinafter diagnostic fragments (DFs).

As used herein, an "isolated nucleic acid molecule" or an "isolated fragment of the *Borrelia burgdorferi* genome" refers to a nucleic acid molecule possessing a specific nucleotide sequence which has been subjected to purification means to reduce, from the composition, the number of compounds which are normally associated with the composition. Particularly, the term refers to the nucleic acid molecules having the sequences set out in SEQ ID NOS: 1-155, to representative fragments thereof as described above including ORF IDs and ORFs, to polynucleotides at least 95%, preferably at least 96%, 97%, 98%, or 99% and especially preferably at least 99.9% identical in sequence thereto, also as set out above.

A variety of purification means can be used to generate the isolated fragments of the present invention. These include, but are not limited to methods which separate constituents of a solution based on charge, solubility, or size.

In one embodiment, *Borrelia burgdorferi* DNA can be enzymatically sheared to produce fragments of 15-20 kb in length. These fragments can then be used to generate a *Borrelia burgdorferi* library by inserting them into lambda clones as described in the Examples below. Primers flanking, for example, an ORF, such as those enumerated in Tables 1-6 can then be generated using nucleotide sequence information provided in SEQ ID NOS: 1-155. Well known and routine techniques of PCR cloning then can be used to isolate the ORF from the lambda DNA library or *Borrelia burgdorferi* genomic DNA. Thus, given the availability of SEQ ID NOS:1-

155, the information in Tables 1-6, and the information that may be obtained readily by analysis of the sequences of SEQ ID NOS:1-155 using methods set out above, those of skill will be enabled by the present disclosure to isolate any ORF-containing or other nucleic acid fragment of the present invention.

The isolated nucleic acid molecules of the present invention include, but are not limited to single stranded and double stranded DNA, and single stranded RNA. For purposes of numbering and reference to polynucleotide and polypeptide sequences the entire sequence of each sequence of SEQ ID NOS:1-155 is included with the first nucleotide being position 1. Therefore, for reference purposes the numbering used in the present invention is that provided in the sequence listing for SEQ ID NOS:1-155.

As used herein, an open reading frame (ORF), means a series of nucleotide triplets coding for amino acid residues without any termination codons and is a sequence translatable into protein. Further, unless specified, the term "ORF" for each ORF ID is defined by the termination codon at the 3' end and the 5' most methionine codon, at the 5' end, in frame with said 3' termination codon. Unless specified, the term "ORF" also refers to a particular polypeptide sequence defined by the ORF polynucleotide sequence, wherein the N-terminus is defined by the 5' most methionine codon in frame with the termination codon at the 3' end of the ORF ID and the C-terminus is defined by the last codon before the said 3' termination codon. As used herein, an ORF ID represents a sequence without any internal termination codons flanked by termination codons.

Tables 1-6 list ORF IDs in the *Borrelia burgdorferi* genomic contigs of the present invention that were identified as putative coding regions by the GeneMark software using organism-specific second-order Markov probability transition matrices. It will be appreciated that other criteria can be used, in accordance with well known analytical methods, such as those discussed herein, to generate more inclusive, more restrictive, or more selective lists.

The *B. burgdorferi* genome consists of one large linear chromosome containing approximately two thirds of its genetic material and multiple extrachromosomal elements (approximately 15) containing the remaining one third of its genetic material. SEQ ID NO:1 (Contig ID 1) is the complete sequence of the large linear *B. burgdorferi* chromosome. SEQ ID NOS:2-155 (Contig ID 2-155 respectively) are fragments (contigs) of the extrachromosomal elements. Tables 1-3 below relate only to SEQ ID NO:1. Tables 4-6 relate to the extrachromosomal elements (SEQ ID NOS:2-155).

Table 1 sets out ORF IDs in the *Borrelia burgdorferi* chromosome of the present invention that cover a continuous region of at least 50 bases are 95% or more identical (by BLAST analysis using default parameters) to a nucleotide sequence available through GenBank in July, 1997.

Table 2 sets out ORF IDs in the *Borrelia burgdorferi* chromosome of the present invention that are not in Table 1 and match, with a BLASTP probability score of 0.01 or less, a polypeptide sequence available through GenBank in July, 1997.

Table 3 sets out ORF IDs in the *Borrelia burgdorferi* chromosome of the present invention that do not match significantly, by BLASTP analysis, a polypeptide sequence available through GenBank in July, 1997.

Table 4 sets out ORF IDs in the *Borrelia burgdorferi* extrachromosomal element contigs of the present invention that over a continuous region of at least 50 bases are 95% or more identical (by BLAST analysis) to a nucleotide sequence available through GenBank in July, 1997.

Table 5 sets out ORF IDs in the *Borrelia burgdorferi* extrachromosomal element contigs of the present invention that are not in Table 1 and match, with a BLASTP probability score of 0.01 or less, a polypeptide sequence available through GenBank in July, 1997.

Table 6 sets out ORF IDs in the *Borrelia burgdorferi* extrachromosomal element contigs of the present invention that do not match significantly, by BLASTP analysis, a polypeptide sequence available through GenBank in July, 1997.

In each table, the first and second columns identify the ORF ID by, respectively, contig number and ORF ID number within the contig; the third column indicates the first nucleotide of the ORF ID, counting from the 5' end of the contig strand; and the fourth column indicates the last nucleotide of the ORF ID, counting from the 5' end of the contig strand.

In Tables 1, 2, 4 and 5, column five, lists the Reference for the closest matching sequence available through GenBank. These reference numbers are the database accession numbers commonly used by those of skill in the art, who will be familiar with their denominators. Descriptions of the nomenclature are available from the National Center for Biotechnology Information. Column seven provides the BLAST identity score from the comparison of the ORF ID and the homologous gene; and column nine indicates the length in nucleotides of the highest scoring segment pair identified by the BLAST identity analysis.

The concepts of percent identity and percent similarity of two polypeptide sequences is well understood in the art. For example, two polypeptides 10 amino acids in length which differ at three amino acid positions (*e.g.*, at positions 1, 3 and 5) are said to have a percent identity of 70%. However, the same two polypeptides would be deemed to have a percent similarity of 80% if, for example at position 5, the amino acids moieties, although not identical, were "similar" (*i.e.*, possessed similar biochemical characteristics). As is known in the art, substitution of one amino acid for a "similar" amino acid is a conservative substitution. Generally, proteins are highly tolerant of conservative substitutions. Many programs for analysis of nucleotide or amino acid sequence similarity, such as fasta and BLAST specifically list percent identity of a matching region as an output parameter. Thus, for instance, Tables 1, 2, 4 and 5 herein enumerate the percent identity and similarity of the highest scoring segment pair in each ORF and its listed relative. Further details concerning the algorithms and criteria used for homology searches are provided below and are described in the pertinent literature highlighted by the citations provided below.

It will be appreciated that other criteria can be used to generate more inclusive and more exclusive listings of the types set out in the tables. As those of skill will appreciate, narrow and broad searches both are useful. Thus, a skilled artisan can readily identify ORFs in contigs of the *Borrelia burgdorferi* genome other than those listed in Tables 1-6, such as ORFs which are overlapping or encoded by the opposite strand of an identified ORF in addition to those ascertainable using the computer-based systems of the present invention.

As used herein, an "expression modulating fragment," EMF, means a series of nucleotide molecules which modulates the expression of an operably linked ORF or EMF.

As used herein, a sequence is said to "modulate the expression of an operably linked sequence" when the expression of the sequence is altered by the presence of the EMF. EMFs include, but are not limited to, promoters, and promoter modulating sequences (inducible elements). One class of EMFs are fragments which induce the expression or an operably linked ORF in response to a specific regulatory factor or physiological event.

EMF sequences can be identified within the contigs of the *Borrelia burgdorferi* genome by their proximity to the ORFs provided in Tables 1-6. An intergenic segment, or a fragment of the intergenic segment, from about 10 to 200 nucleotides in length, taken from any one of the ORFs of Tables 1-6 will modulate the expression of an operably linked ORF in a fashion similar to that found with the naturally linked ORF sequence. As used herein, an "intergenic segment" refers to fragments of the *Borrelia burgdorferi* genome which are between two ORF(s) herein described. EMFs also can be identified using known EMFs as a target sequence or target motif in the computer-based systems of the present invention. Further, the two methods can be combined and used together.

The presence and activity of an EMF can be confirmed using an EMF trap vector. An EMF trap vector contains a cloning site linked to a marker sequence. A marker sequence encodes an identifiable phenotype, such as antibiotic resistance or a complementing nutrition auxotrophic factor, which can be identified or assayed when the EMF trap vector is placed within an appropriate host under appropriate conditions. As described above, a EMF will modulate the expression of an operably linked marker sequence. A more detailed discussion of various marker sequences is provided below. A sequence which is suspected as being an EMF is cloned in all three reading frames in one or more restriction sites upstream from the marker sequence in the EMF trap vector. The vector is then transformed into an appropriate host using known procedures and the phenotype of the transformed host in examined under appropriate conditions. As described above, an EMF will modulate the expression of an operably linked marker sequence.

As used herein, a "diagnostic fragment," DF, means a series of nucleotide molecules which selectively hybridize to *Borrelia burgdorferi* sequences. DFs can be readily identified by identifying unique sequences within contigs of the *Borrelia burgdorferi* genome, such as by using well-known computer analysis software, and by generating and testing probes or

amplification primers consisting of the DF sequence in an appropriate diagnostic format which determines amplification or hybridization selectivity.

The sequences falling within the scope of the present invention are not limited to the specific sequences herein described, but also include allelic and species variations thereof. Allelic and species variations can be routinely determined by comparing the sequences provided in SEQ ID NOS:1-155, ORF IDs and ORFs within, a representative fragment thereof, or a nucleotide sequence at least 99% and preferably 99.9% identical to SEQ ID NOS: 1-155, ORF IDs and ORFs within, with a sequence from another isolate of the same species. Furthermore, to accommodate codon variability, the invention includes nucleic acid molecules coding for the same amino acid sequences as do the specific ORFs disclosed herein. In other words, in the coding region of an ORF, substitution of one codon for another which encodes the same amino acid is expressly contemplated.

Any specific sequence disclosed herein can be readily screened for errors by resequencing a particular fragment, such as an ORF, in both directions (*i.e.*, sequence both strands). Alternatively, error screening can be performed by sequencing corresponding polynucleotides of *Borrelia burgdorferi* origin isolated by using part or all of the fragments in question as a probe or primer.

Each of the ORF IDs and ORFs of the *Borrelia burgdorferi* genome disclosed in Tables 1-6, and the EMFs found 5' to the ORF IDs, can be used as polynucleotide reagents in numerous ways. For example, the sequences can be used as diagnostic probes or diagnostic amplification primers to detect the presence of a specific microbe in a sample, particularly *Borrelia burgdorferi*. Especially preferred in this regard are ORF IDs and ORFs such as those of Tables 3 and 6, which do not match previously characterized sequences from other organisms and thus are most likely to be highly selective for *Borrelia burgdorferi*. Also particularly preferred are ORF IDs and ORFs that can be used to distinguish between strains of *Borrelia burgdorferi*, particularly those that distinguish medically important strain, such as drug-resistant strains.

In addition, the fragments of the present invention, as broadly described, can be used to control gene expression through triple helix formation or antisense DNA or RNA, both of which methods are based on the binding of a polynucleotide sequence to DNA or RNA. Triple helix-formation optimally results in a shut-off of RNA transcription from DNA, while antisense RNA hybridization blocks translation of an mRNA molecule into polypeptide. Information from the sequences of the present invention can be used to design antisense and triple helix-forming oligonucleotides. Polynucleotides suitable for use in these methods are usually 20 to 40 bases in length and are designed to be complementary to a region of the gene involved in transcription, for triple-helix formation, or to the mRNA itself, for antisense inhibition. Both techniques have been demonstrated to be effective in model systems, and the requisite techniques are well known and involve routine procedures. Triple helix techniques are discussed in, for example, Lee *et al.*, *Nucl. Acids Res. 6*:3073 (1979); Cooney *et al.*, *Science 241*:456 (1988); and Dervan *et al.*, *Science 251*:1360 (1991). Antisense techniques in general are discussed in, for instance, Okano,

J. Neurochem. *56*:560 (1991) and *Oligodeoxynucleotides as Antisense Inhibitors of Gene Expression*, CRC Press, Boca Raton, FL (1988)).

The present invention further provides recombinant constructs comprising one or more fragments of the *Borrelia burgdorferi* genomic fragments and contigs of the present invention. Certain preferred recombinant constructs of the present invention comprise a vector, such as a plasmid or viral vector, into which a fragment of the *Borrelia burgdorferi* genome has been inserted, in a forward or reverse orientation. In the case of a vector comprising one of the ORF IDs or ORFs of the present invention, the vector may further comprise regulatory sequences, including for example, a promoter, operably linked to the ORF ID or ORF. For vectors comprising the EMFs of the present invention, the vector may further comprise a marker sequence or heterologous ORF ID or ORF operably linked to the EMF.

Large numbers of suitable vectors and promoters are known to those of skill in the art and are commercially available for generating the recombinant constructs of the present invention. The following vectors are provided by way of example. Useful bacterial vectors include phagescript, PsiX174, pBluescript SK, pBS KS, pNH8a, pNH16a, pNH18a, pNH46a (available from Stratagene); pTrc99A, pKK223-3, pKK233-3, pDR540, pRIT5 (available from Pharmacia); pQE vectors (available from Promega). Useful eukaryotic vectors include pWLneo, pSV2cat, pOG44, pXT1, pSG (available from Stratagene) pSVK3, pBPV, pMSG, pSVL (available from Pharmacia).

Promoter regions can be selected from any desired gene using CAT (chloramphenicol transferase) vectors or other vectors with selectable markers. Two appropriate vectors are pKK232-8 and pCM7. Particular named bacterial promoters include lacI, lacZ, T3, T7, gpt, lambda PR, and trc. Eukaryotic promoters include CMV immediate early, HSV thymidine kinase, early and late SV40, LTRs from retrovirus, and mouse metallothionein- I. Selection of the appropriate vector and promoter is well within the level of ordinary skill in the art.

The present invention further provides host cells containing any one of the isolated fragments of the *Borrelia burgdorferi* genomic fragments and contigs of the present invention, wherein the fragment has been introduced into the host cell using known methods. The host cell can be a higher eukaryotic host cell, such as a mammalian cell, a lower eukaryotic host cell, such as a yeast cell, or a procaryotic cell, such as a bacterial cell.

A polynucleotide of the present invention, such as a recombinant construct comprising an ORF of the present invention, may be introduced into the host by a variety of well established techniques that are standard in the art, such as calcium phosphate transfection, DEAE, dextran mediated transfection and electroporation, which are described in, for instance, Davis, L. *et al.*, BASIC METHODS IN MOLECULAR BIOLOGY (1986).

A host cell containing one of the fragments of the *Borrelia burgdorferi* genomic fragments and contigs of the present invention, can be used in conventional manners to produce the gene product encoded by the isolated fragment (in the case of an ORF) or can be used to produce a heterologous protein under the control of the EMF.

The present invention further provides isolated polypeptides encoded by the nucleic acid fragments of the present invention or by degenerate variants of the nucleic acid fragments of the present invention. By "degenerate variant" is intended nucleotide fragments which differ from a nucleic acid fragment of the present invention (*e.g.*, an ORF) by nucleotide sequence but, due to the degeneracy of the Genetic Code, encode an identical polypeptide sequence.

Preferred nucleic acid fragments of the present invention are the ORF IDs depicted in Tables 2, 3, 5 and 6, and ORFs witin, which encode proteins.

A variety of methodologies known in the art can be utilized to obtain any one of the isolated polypeptides or proteins of the present invention. At the simplest level, the amino acid sequence can be synthesized using commercially available peptide synthesizers. This is particularly useful in producing small peptides and fragments of larger polypeptides. Such short fragments as may be obtained most readily by synthesis are useful, for example, in generating antibodies against the native polypeptide, as discussed further below.

In an alternative method, the polypeptide or protein is purified from bacterial cells which naturally produce the polypeptide or protein. One skilled in the art can readily employ well-known methods for isolating polypeptides and proteins to isolate and purify polypeptides or proteins of the present invention produced naturally by a bacterial strain, or by other methods. Methods for isolation and purification that can be employed in this regard include, but are not limited to, immunochromatography, HPLC, size-exclusion chromatography, ion-exchange chromatography, and immuno-affinity chromatography.

The polypeptides and proteins of the present invention also can be purified from cells which have been altered to express the desired polypeptide or protein. As used herein, a cell is said to be altered to express a desired polypeptide or protein when the cell, through genetic manipulation, is made to produce a polypeptide or protein which it normally does not produce or which the cell normally produces at a lower level. Those skilled in the art can readily adapt procedures for introducing and expressing either recombinant or synthetic sequences into eukaryotic or prokaryotic cells in order to generate a cell which produces one of the polypeptides or proteins of the present invention.

The polypeptides of the present invention are preferably provided in an isolated form, and preferably are substantially purified. A recombinantly produced version of the *B. burgdorferi* polypeptide can be substantially purified by the one-step method described by Smith et al. (1988) Gene 67:31-40. Polypeptides of the invention also can be purified from natural or recombinant sources using antibodies directed against the polypeptides of the invention in methods which are well known in the art of protein purification.

The invention further provides for isolated *B. burgdorferi* polypeptides comprising an amino acid sequence selected from the group including: (a) the amino acid sequence of a full-length *B. burgdorferi* polypeptide having the complete amino acid sequence from the first methionine codon to the termination codon of each sequence listed in SEQ ID NOS:1-155, wherein said termination codon is at the end of each SEQ ID NO: and said first methionine is the

first methionine in frame with said termination codon; and (b) the amino acid sequence of a full-length *B. burgdorferi* polypeptide having the complete amino acid sequence in (a) excepting the N-terminal methionine.

The polypeptides of the present invention also include polypeptides having an amino acid sequence at least 80% identical, more preferably at least 90% identical, and still more preferably 95%, 96%, 97%, 98% or 99% identical to those described in (a) and (b) above.

The present invention is further directed to polynucleotides encoding portions or fragments of the amino acid sequences described herein as well as to portions or fragments of the isolated amino acid sequences described herein. Fragments include portions of the amino acid sequences described herein at least 5 contiguous amino acid in length and selected from any two integers, one of which representing an N-terminal position and another representing a C-terminal position. The initiation codon of the ORFs of the present invention is position 1. The initiation codon (positon 1) for purposes of the present invention is the first methionine codon of each ORF ID which is in frame with the termination codon at the end of each said sequence. Every combination of a N-terminal and C-terminal position that a fragment at least 5 contiguous amino acid residues in length could occupy, on any given ORF is included in the invention, i.e., from initiation codon up to the termination codon. "At least" means a fragment may be 5 contiguous amino acid residues in length or any integer between 5 and the number of residues in an ORF, minus 1. Therefore, included in the invention are contiguous fragments specified by any N-terminal and C-terminal positions of amino acid sequence set forth in SEQ ID NOS:1-155 or Tables 1-6 wherein the contiguous fragment is any integer between 5 and the number of residues in an ORF minus 1.

Further, the invention includes polypeptides comprising fragments specified by size, in amino acid residues, rather than by N-terminal and C-terminal positions. The invention includes any fragment size, in contiguous amino acid residues, selected from integers between 5 and the number of residues in an ORF, minus 1. Preferred sizes of contiguous polypeptide fragments include about 5 amino acid residues, about 10 amino acid residues, about 20 amino acid residues, about 30 amino acid residues, about 40 amino acid residues, about 50 amino acid residues, about 100 amino acid residues, about 200 amino acid residues, about 300 amino acid residues, and about 400 amino acid residues. The preferred sizes are, of course, meant to exemplify, not limit, the present invention as all size fragments representing any integer between 5 and the number of residues in a full length sequence minus 1 are included in the invention. The present invention also provides for the exclusion of any fragments specified by N-terminal and C-terminal positions or by size in amino acid residues as described above. Any number of fragments specified by N-terminal and C-terminal positions or by size in amino acid residues as described above may be excluded.

The above fragments need not be active since they would be useful, for example, in immunoassays, in epitope mapping, epitope tagging, to generate antibodies to a particular portion of the protein, as vaccines, and as molecular weight markers.

Further polypeptides of the present invention include polypeptides which have at least 90% similarity, more preferably at least 95% similarity, and still more preferably at least 96%, 97%, 98% or 99% similarity to those described above.

A further embodiment of the invention relates to a polypeptide which comprises the amino acid sequence of a *B. burgdorferi* polypeptide having an amino acid sequence which contains at least one conservative amino acid substitution, but not more than 50 conservative amino acid substitutions, not more than 40 conservative amino acid substitutions, not more than 30 conservative amino acid substitutions, and not more than 20 conservative amino acid substitutions. Also provided are polypeptides which comprise the amino acid sequence of a *B. burgdorferi* polypeptide, having at least one, but not more than 10, 9, 8, 7, 6, 5, 4, 3, 2 or 1 conservative amino acid substitutions.

By a polypeptide having an amino acid sequence at least, for example, 95% "identical" to a query amino acid sequence of the present invention, it is intended that the amino acid sequence of the subject polypeptide is identical to the query sequence except that the subject polypeptide sequence may include up to five amino acid alterations per each 100 amino acids of the query amino acid sequence. In other words, to obtain a polypeptide having an amino acid sequence at least 95% identical to a query amino acid sequence, up to 5% of the amino acid residues in the subject sequence may be inserted, deleted, (indels) or substituted with another amino acid. These alterations of the reference sequence may occur at the amino or carboxy terminal positions of the reference amino acid sequence or anywhere between those terminal positions, interspersed either individually among residues in the reference sequence or in one or more contiguous groups within the reference sequence.

As a practical matter, whether any particular polypeptide is at least 90%, 95%, 96%, 97%, 98% or 99% identical to the ORF amino acid sequences encoded by the sequences of SEQ ID NOS:1-155, as described hererin, can be determined conventionally using known computer programs. A preferred method for determining the best overall match between a query sequence (a sequence of the present invention) and a subject sequence, also referred to as a global sequence alignment, can be determined using the FASTDB computer program based on the algorithm of Brutlag et al., (1990) Comp. App. Biosci. 6:237-245. In a sequence alignment the query and subject sequences are both amino acid sequences. The result of said global sequence alignment is in percent identity. Preferred parameters used in a FASTDB amino acid alignment are: Matrix=PAM 0, k-tuple=2, Mismatch Penalty=1, Joining Penalty=20, Randomization Group Length=0, Cutoff Score=1, Window Size=sequence length, Gap Penalty=5, Gap Size Penalty=0.05, Window Size=500 or the length of the subject amino acid sequence, whichever is shorter.

If the subject sequence is shorter than the query sequence due to N- or C-terminal deletions, not because of internal deletions, the results, in percent identity, must be manually corrected. This is because the FASTDB program does not account for N- and C-terminal truncations of the subject sequence when calculating global percent identity. For subject

sequences truncated at the N- and C-termini, relative to the query sequence, the percent identity is corrected by calculating the number of residues of the query sequence that are N- and C-terminal of the subject sequence, which are not matched/aligned with a corresponding subject residue, as a percent of the total bases of the query sequence. Whether a residue is matched/aligned is determined by results of the FASTDB sequence alignment. This percentage is then subtracted from the percent identity, calculated by the above FASTDB program using the specified parameters, to arrive at a final percent identity score. This final percent identity score is what is used for the purposes of the present invention. Only residues to the N- and C-termini of the subject sequence, which are not matched/aligned with the query sequence, are considered for the purposes of manually adjusting the percent identity score. That is, only query amino acid residues outside the farthest N- and C-terminal residues of the subject sequence.

For example, a 90 amino acid residue subject sequence is aligned with a 100 residue query sequence to determine percent identity. The deletion occurs at the N-terminus of the subject sequence and therefore, the FASTDB alignment does not match/align with the first 10 residues at the N-terminus. The 10 unpaired residues represent 10% of the sequence (number of residues at the N- and C- termini not matched/total number of residues in the query sequence) so 10% is subtracted from the percent identity score calculated by the FASTDB program. If the remaining 90 residues were perfectly matched the final percent identity would be 90%. In another example, a 90 residue subject sequence is compared with a 100 residue query sequence. This time the deletions are internal so there are no residues at the N- or C-termini of the subject sequence which are not matched/aligned with the query. In this case the percent identity calculated by FASTDB is not manually corrected. Once again, only residue positions outside the N- and C-terminal ends of the subject sequence, as displayed in the FASTDB alignment, which are not matched/aligned with the query sequence are manually corrected. No other manual corrections are to made for the purposes of the present invention.

The above polypeptide sequences are included irrespective of whether they have their normal biological activity. This is because even where a particular polypeptide molecule does not have biological activity, one of skill in the art would still know how to use the polypeptide, for instance, as a vaccine or to generate antibodies. Other uses of the polypeptides of the present invention that do not have *B. burgdorferi* activity include, *inter alia*, as epitope tags, in epitope mapping, and as molecular weight markers on SDS-PAGE gels or on molecular sieve gel filtration columns using methods known to those of skill in the art.

As described below, the polypeptides of the present invention can also be used to raise polyclonal and monoclonal antibodies, which are useful in assays for detecting *B. burgdorferi* protein expression or as agonists and antagonists capable of enhancing or inhibiting *B. burgdorferi* protein function. Further, such polypeptides can be used in the yeast two-hybrid system to "capture" *B. burgdorferi* protein binding proteins which are also candidate agonists and antagonists according to the present invention. *See, e.g.,* Fields et al. (1989) Nature 340:245-246.

Any host/vector system can be used to express one or more of the ORFs of the present invention. These include, but are not limited to, eukaryotic hosts such as HeLa cells, CV-1 cell, COS cells, and Sf9 cells, as well as prokaryotic host such as *E. coli* and *B. subtilis*. The most preferred cells are those which do not normally express the particular polypeptide or protein or which expresses the polypeptide or protein at low natural level.

"Recombinant," as used herein, means that a polypeptide or protein is derived from recombinant (*e.g.*, microbial or mammalian) expression systems. "Microbial" refers to recombinant polypeptides or proteins made in bacterial or fungal (*e.g.*, yeast) expression systems. As a product, "recombinant microbial"defines a polypeptide or protein essentially free of native endogenous substances and unaccompanied by associated native glycosylation. Polypeptides or proteins expressed in most bacterial cultures, *e.g.*, *E. coli*, will be free of glycosylation modifications; polypeptides or proteins expressed in yeast will have a glycosylation pattern different from that expressed in mammalian cells.

"Nucleotide sequence" refers to a heteropolymer of deoxyribonucleotides. Generally, DNA segments encoding the polypeptides and proteins provided by this invention are assembled from fragments of the *Borrelia burgdorferi* genome and short oligonucleotide linkers, or from a series of oligonucleotides, to provide a synthetic gene which is capable of being expressed in a recombinant transcriptional unit comprising regulatory elements derived from a microbial or viral operon.

Recombinant expression vehicle or vector" refers to a plasmid or phage or virus or vector, for expressing a polypeptide from a DNA (RNA) sequence. The expression vehicle can comprise a transcriptional unit comprising an assembly of (1) a genetic regulatory elements necessary for gene expression in the host, including elements required to initiate and maintain transcription at a level sufficient for suitable expression of the desired polypeptide, including, for example, promoters and, where necessary, an enhancer and a polyadenylation signal; (2) a structural or coding sequence which is transcribed into mRNA and translated into protein, and (3) appropriate signals to initiate translation at the beginning of the desired coding region and terminate translation at its end. Structural units intended for use in yeast or eukaryotic expression systems preferably include a leader sequence enabling extracellular secretion of translated protein by a host cell. Alternatively, where recombinant protein is expressed without a leader or transport sequence, it may include an N-terminal methionine residue. This residue may or may not be subsequently cleaved from the expressed recombinant protein to provide a final product.

"Recombinant expression system" means host cells which have stably integrated a recombinant transcriptional unit into chromosomal DNA or carry the recombinant transcriptional unit extra chromosomally. The cells can be prokaryotic or eukaryotic. Recombinant expression systems as defined herein will express heterologous polypeptides or proteins upon induction of the regulatory elements linked to the DNA segment or synthetic gene to be expressed.

Mature proteins can be expressed in mammalian cells, yeast, bacteria, or other cells under the control of appropriate promoters. Cell-free translation systems can also be employed to

produce such proteins using RNAs derived from the DNA constructs of the present invention. Appropriate cloning and expression vectors for use with prokaryotic and eukaryotic hosts are described in Sambrook *et al.*, *Molecular Cloning*: *A Laboratory Manual,* 2nd Edition, Cold Spring Harbor Laboratory Press, Cold Spring Harbor, New York (1989), the disclosure of which is hereby incorporated by reference in its entirety.

Generally, recombinant expression vectors will include origins of replication and selectable markers permitting transformation of the host cell, *e.g.*, the ampicillin resistance gene of *E. coli* and *S. cerevisiae* TRP1 gene, and a promoter derived from a highly expressed gene to direct transcription of a downstream structural sequence. Such promoters can be derived from operons encoding glycolytic enzymes such as 3- phosphoglycerate kinase (PGK), alpha-factor, acid phosphatase, or heat shock proteins, among others. The heterologous structural sequence is assembled in appropriate phase with translation initiation and termination sequences, and preferably, a leader sequence capable of directing secretion of translated protein into the periplasmic space or extracellular medium. Optionally, the heterologous sequence can encode a fusion protein including an N-terminal identification peptide imparting desired characteristics, e.g., stabilization or simplified purification of expressed recombinant product.

Useful expression vectors for bacterial use are constructed by inserting a structural DNA sequence encoding a desired protein together with suitable translation initiation and termination signals in operable reading phase with a functional promoter. The vector will comprise one or more phenotypic selectable markers and an origin of replication to ensure maintenance of the vector and, when desirable, provide amplification within the host.

Suitable prokaryotic hosts for transformation include strains of *E. coli, B. subtilis, Salmonella typhimurium* and various species within the genera *Pseudomonas* and *Streptomyces*. Others may, also be employed as a matter of choice.

As a representative but non-limiting example, useful expression vectors for bacterial use can comprise a selectable marker and bacterial origin of replication derived from commercially available plasmids comprising genetic elements of the well known cloning vector pBR322 (ATCC 37017). Such commercial vectors include, for example, pKK223-3 (available form Pharmacia Fine Chemicals, Uppsala, Sweden) and GEM 1 (available from Promega Biotec, Madison, WI, USA). These pBR322 "backbone" sections are combined with an appropriate promoter and the structural sequence to be expressed.

Following transformation of a suitable host strain and growth of the host strain to an appropriate cell density, the selected promoter, where it is inducible, is derepressed or induced by appropriate means (*e.g.*, temperature shift or chemical induction) and cells are cultured for an additional period to provide for expression of the induced gene product. Thereafter cells are typically harvested, generally by centrifugation, disrupted to release expressed protein, generally by physical or chemical means, and the resulting crude extract is retained for further purification.

Various mammalian cell culture systems can also be employed to express recombinant protein. Examples of mammalian expression systems include the COS-7 lines of monkey kidney

fibroblasts, described in Gluzman, *Cell 23*:175 (1981), and other cell lines capable of expressing a compatible vector, for example, the C127, 3T3, CHO, HeLa and BHK cell lines.

Mammalian expression vectors will comprise an origin of replication, a suitable promoter and enhancer, and also any necessary ribosome binding sites, polyadenylation site, splice donor and acceptor sites, transcriptional termination sequences, and 5' flanking nontranscribed sequences. DNA sequences derived from the SV40 viral genome, for example, SV40 origin, early promoter, enhancer, splice, and polyadenylation sites may be used to provide the required nontranscribed genetic elements.

Recombinant polypeptides and proteins produced in bacterial culture is usually isolated by initial extraction from cell pellets, followed by one or more salting-out, aqueous ion exchange or size exclusion chromatography steps. Microbial cells employed in expression of proteins can be disrupted by any convenient method, including freeze-thaw cycling, sonication, mechanical disruption, or use of cell lysing agents. Protein refolding steps can be used, as necessary, in completing configuration of the mature protein. Finally, high performance liquid chromatography (HPLC) can be employed for final purification steps.

The present invention further includes isolated polypeptides, proteins and nucleic acid molecules which are substantially equivalent to those herein described. As used herein, substantially equivalent can refer both to nucleic acid and amino acid sequences, for example a mutant sequence, that varies from a reference sequence by one or more substitutions, deletions, or additions, the net effect of which does not result in an adverse functional dissimilarity between reference and subject sequences. Particularly preferred in this regard are conservative substitutions, known to those of skill in the art. For purposes of the present invention, sequences having equivalent biological activity, and equivalent expression characteristics are considered substantially equivalent. For purposes of determining equivalence, truncation of the mature sequence (e.g., removal of leader sequence(s)) should be disregarded.

The invention further provides methods of obtaining homologs from other strains of *Borrelia burgdorferi*, of the fragments of the *Borrelia burgdorferi* genome of the present invention and homologs of the proteins encoded by the ORFs of the present invention. As used herein, a sequence or protein of *Borrelia burgdorferi* is defined as a homolog of a fragment of the *Borrelia burgdorferi* fragments or contigs or a protein encoded by one of the ORFs of the present invention, if it shares significant homology to one of the fragments of the *Borrelia burgdorferi* genome of the present invention or a protein encoded by one of the ORFs of the present invention. Specifically, by using the sequence disclosed herein as a probe or as primers, and techniques such as PCR cloning and colony/plaque hybridization, one skilled in the art can obtain homologs.

As used herein, two nucleic acid molecules or proteins are said to "share significant homology" if the two contain regions which possess greater than 85% sequence (amino acid or nucleic acid) homology. Preferred homologs in this regard are those with more than 90% homology. Especially preferred are those with 95% or more homology. Among especially

preferred homologs those with 96, 97%, 98%, 99% or more homology are particularly preferred. The most preferred homologs among these are those with 99.9% homology or more. It will be understood that, among measures of homology, identity is particularly preferred in this regard.

Region specific primers or probes derived from the nucleotide sequence provided in SEQ ID NOS: 1-155 or from a nucleotide sequence at least 95%, particularly at least 96%, 97%, 98% or 99%, especially at least 99.5% identical to a sequence of SEQ ID NOS: 1-155 can be used to prime DNA synthesis and PCR amplification, as well as to identify colonies containing cloned DNA encoding a homolog. Methods suitable to this aspect of the present invention are well known and have been described in great detail in many publications such as, for example, Innis *et al.*, *PCR Protocols*, Academic Press, San Diego, CA (1990)).

When using primers derived from SEQ ID NOS: 1-155 or from a nucleotide sequence having an aforementioned identity to a sequence of SEQ ID NOS:1-155, one skilled in the art will recognize that by employing high stringency conditions (*e.g.*, annealing at 50-60°C in 6X SSPC and 50% formamide, and washing at 50- 65°C in 0.5X SSPC) only sequences which are greater than 75% homologous to the primer will be amplified. By employing lower stringency conditions (*e.g.*, hybridizing at 35-37°C in 5X SSPC and 40-45% formamide, and washing at 42°C in 0.5X SSPC), sequences which are greater than 40-50% homologous to the primer will also be amplified.

When using DNA probes derived from SEQ ID NOS:1-155, or from a nucleotide sequence having an aforementioned identity to a sequence of SEQ ID NOS: 1-155 , for colony/plaque hybridization, one skilled in the art will recognize that by employing high stringency conditions (*e.g.*, hybridizing at 50- 65°C in 5X SSPC and 50% formamide, and washing at 50- 65°C in 0.5X SSPC), sequences having regions which are greater than 90% homologous to the probe can be obtained, and that by employing lower stringency conditions (*e.g.*, hybridizing at 35-37°C in 5X SSPC and 40-45% formamide, and washing at 42°C in 0.5X SSPC), sequences having regions which are greater than 35-45% homologous to the probe will be obtained.

Any organism can be used as the source for homologs of the present invention so long as the organism naturally expresses such a protein or contains genes encoding the same. The most preferred organism for isolating homologs are bacteria which are closely related to *Borrelia burgdorferi*.

ILLUSTRATIVE USES OF COMPOSITIONS OF THE INVENTION

Each ORF of the ORF IDs provided in Tables 1, 2, 4 and 5 is identified with a function by homology to a known gene or polypeptide. As a result, one skilled in the art can use the polypeptides of the present invention for commercial, therapeutic and industrial purposes consistent with the type of putative identification of the polypeptide. Such identifications permit one skilled in the art to use the *Borrelia burgdorferi* ORFs in a manner similar to the known type

of sequences for which the identification is made; for example, to ferment a particular sugar source or to produce a particular metabolite. A variety of reviews illustrative of this aspect of the invention are available, including the following reviews on the industrial use of enzymes, for example, BIOCHEMICAL ENGINEERING AND BIOTECHNOLOGY HANDBOOK, 2nd Ed., MacMillan Publications, Ltd. NY (1991) and BIOCATALYSTS IN ORGANIC SYNTHESES, Tramper *et al.*, Eds., Elsevier Science Publishers, Amsterdam, The Netherlands (1985). A variety of exemplary uses that illustrate this and similar aspects of the present invention are discussed below.

1. Biosynthetic Enzymes

Open reading frames encoding proteins involved in mediating the catalytic reactions involved in intermediary and macromolecular metabolism, the biosynthesis of small molecules, cellular processes and other functions includes enzymes involved in the degradation of the intermediary products of metabolism, enzymes involved in central intermediary metabolism, enzymes involved in respiration, both aerobic and anaerobic, enzymes involved in fermentation, enzymes involved in ATP proton motor force conversion, enzymes involved in broad regulatory function, enzymes involved in amino acid synthesis, enzymes involved in nucleotide synthesis, enzymes involved in cofactor and vitamin synthesis, can be used for industrial biosynthesis.

The various metabolic pathways present in *Borrelia burgdorferi* can be identified based on absolute nutritional requirements as well as by examining the various enzymes identified in Table 1-6 and SEQ ID NOS:1-155.

Of particular interest are polypeptides involved in the degradation of intermediary metabolites as well as non-macromolecular metabolism. Such enzymes include amylases, glucose oxidases, and catalase.

Proteolytic enzymes are another class of commercially important enzymes. Proteolytic enzymes find use in a number of industrial processes including the processing of flax and other vegetable fibers, in the extraction, clarification and depectinization of fruit juices, in the extraction of vegetables' oil and in the maceration of fruits and vegetables to give unicellular fruits. A detailed review of the proteolytic enzymes used in the food industry is provided in Rombouts *et al.*, *Symbiosis 21:*79 (1986) and Voragen *et al.* in *Biocatalysts In Agricultural Biotechnology*, Whitaker *et al.*, Eds., *American Chemical Society Symposium Series 389:*93 (1989) .

The metabolism of sugars is an important aspect of the primary metabolism of *Borrelia burgdorferi*. Enzymes involved in the degradation of sugars, such as, particularly, glucose, galactose, fructose and xylose, can be used in industrial fermentation. Some of the important sugar transforming enzymes, from a commercial viewpoint, include sugar isomerases such as glucose isomerase. Other metabolic enzymes have found commercial use such as glucose oxidases which produces ketogulonic acid (KGA). KGA is an intermediate in the commercial production of ascorbic acid using the Reichstein's procedure, as described in Krueger *et al.*, *Biotechnology* 6(A), Rhine *et al.*, Eds., Verlag Press, Weinheim, Germany (1984).

Glucose oxidase (GOD) is commercially available and has been used in purified form as well as in an immobilized form for the deoxygenation of beer. See, for instance, Hartmeir *et al.*, *Biotechnology Letters 1*:21 (1979). The most important application of GOD is the industrial scale fermentation of gluconic acid. Market for gluconic acids which are used in the detergent, textile, leather, photographic, pharmaceutical, food, feed and concrete industry, as described, for example, in Bigelis *et al.*, beginning on page 357 in GENE MANIPULATIONS AND FUNGI; Benett *et al.*, Eds., Academic Press, New York (1985). In addition to industrial applications, GOD has found applications in medicine for quantitative determination of glucose in body fluids recently in biotechnology for analyzing syrups from starch and cellulose hydrosylates. This application is described in Owusu *et al.*, *Biochem. et Biophysica. Acta. 872:*83 (1986), for instance.

The main sweetener used in the world today is sugar which comes from sugar beets and sugar cane. In the field of industrial enzymes, the glucose isomerase process shows the largest expansion in the market today. Initially, soluble enzymes were used and later immobilized enzymes were developed (Krueger *et al.*, *Biotechnology, The Textbook of Industrial Microbiology*, Sinauer Associated Incorporated, Sunderland, Massachusetts (1990)). Today, the use of glucose- produced high fructose syrups is by far the largest industrial business using immobilized enzymes. A review of the industrial use of these enzymes is provided by Jorgensen, Starch *40*:307 (1988).

Proteinases, such as alkaline serine proteinases, are used as detergent additives and thus represent one of the largest volumes of microbial enzymes used in the industrial sector. Because of their industrial importance, there is a large body of published and unpublished information regarding the use of these enzymes in industrial processes. (See Faultman *et al.*, Acid Proteases Structure Function and Biology, Tang, J., ed., Plenum Press, New York (1977) and Godfrey *et al.*, Industrial Enzymes, MacMillan Publishers, Surrey, UK (1983) and Hepner *et al.*, Report Industrial Enzymes by 1990, Hel Hepner & Associates, London (1986)).

Another class of commercially usable proteins of the present invention are the microbial lipases, described by, for instance, Macrae *et al.*, *Philosophical Transactions of the Chiral Society of London 310*:227 (1985) and Poserke, *Journal of the American Oil Chemist Society 61:*1758 (1984). A major use of lipases is in the fat and oil industry for the production of neutral glycerides using lipase catalyzed inter-esterification of readily available triglycerides. Application of lipases include the use as a detergent additive to facilitate the removal of fats from fabrics in the course of the washing procedures.

The use of enzymes, and in particular microbial enzymes, as catalyst for key steps in the synthesis of complex organic molecules is gaining popularity at a great rate. One area of great interest is the preparation of chiral intermediates. Preparation of chiral intermediates is of interest to a wide range of synthetic chemists particularly those scientists involved with the preparation of new pharmaceuticals, agrochemicals, fragrances and flavors. (See Davies *et al.*, *Recent Advances in the Generation of Chiral Intermediates Using Enzymes*, CRC Press, Boca Raton,

Florida (1990)). The following reactions catalyzed by enzymes are of interest to organic chemists: hydrolysis of carboxylic acid esters, phosphate esters, amides and nitriles, esterification reactions, trans-esterification reactions, synthesis of amides, reduction of alkanones and oxoalkanates, oxidation of alcohols to carbonyl compounds, oxidation of sulfides to sulfoxides, and carbon bond forming reactions such as the aldol reaction.

When considering the use of an enzyme encoded by one of the ORFs of the present invention for biotransformation and organic synthesis it is sometimes necessary to consider the respective advantages and disadvantages of using a microorganism as opposed to an isolated enzyme. Pros and cons of using a whole cell system on the one hand or an isolated partially purified enzyme on the other hand, has been described in detail by Bud *et al.*, Chemistry in Britain (1987), p. 127.

Amino transferases, enzymes involved in the biosynthesis and metabolism of amino acids, are useful in the catalytic production of amino acids. The advantages of using microbial based enzyme systems is that the amino transferase enzymes catalyze the stereo- selective synthesis of only L-amino acids and generally possess uniformly high catalytic rates. A description of the use of amino transferases for amino acid production is provided by Roselle-David, *Methods of Enzymology 136*:479 (1987).

Another category of useful proteins encoded by the ORFs of the present invention include enzymes involved in nucleic acid synthesis, repair, and recombination.

2. Generation of Antibodies

As described here, the proteins of the present invention, as well as homologs thereof, can be used in a variety of procedures and methods known in the art which are currently applied to other proteins. The proteins of the present invention can further be used to generate an antibody which selectively binds the protein.

B. burgdorferi protein-specific antibodies for use in the present invention can be raised against the intact *B. burgdorferi* protein or an antigenic polypeptide fragment thereof, which may be presented together with a carrier protein, such as an albumin, to an animal system (such as rabbit or mouse) or, if it is long enough (at least about 25 amino acids), without a carrier.

As used herein, the term "antibody" (Ab) or "monoclonal antibody" (Mab) is meant to include intact molecules, single chain whole antibodies, and antibody fragments. Antibody fragments of the present invention include Fab and F(ab')2 and other fragments including single-chain Fvs (scFv) and disulfide-linked Fvs (sdFv). Also included in the present invention are chimeric and humanized monoclonal antibodies and polyclonal antibodies specific for the polypeptides of the present invention. The antibodies of the present invention may be prepared by any of a variety of methods. For example, cells expressing a polypeptide of the present invention or an antigenic fragment thereof can be administered to an animal in order to induce the production of sera containing polyclonal antibodies. For example, a preparation of *B. burgdorferi* polypeptide or fragment thereof is prepared and purified to render it substantially free

of natural contaminants. Such a preparation is then introduced into an animal in order to produce polyclonal antisera of greater specific activity.

In a preferred method, the antibodies of the present invention are monoclonal antibodies or binding fragments thereof. Such monoclonal antibodies can be prepared using hybridoma technology. *See, e.g.,* Harlow et al., ANTIBODIES: A LABORATORY MANUAL, (Cold Spring Harbor Laboratory Press, 2nd ed. 1988); Hammerling, et al., in: MONOCLONAL ANTIBODIES AND T-CELL HYBRIDOMAS 563-681 (Elsevier, N.Y., 1981). Fab and F(ab')2 fragments may be produced by proteolytic cleavage, using enzymes such as papain (to produce Fab fragments) or pepsin (to produce F(ab')2 fragments). Alternatively, *B. burgdorferi* polypeptide-binding fragments, chimeric, and humanized antibodies can be produced through the application of recombinant DNA technology or through synthetic chemistry using methods known in the art.

Alternatively, additional antibodies capable of binding to the polypeptide antigen of the present invention may be produced in a two-step procedure through the use of anti-idiotypic antibodies. Such a method makes use of the fact that antibodies are themselves antigens, and that, therefore, it is possible to obtain an antibody which binds to a second antibody. In accordance with this method, *B. burgdorferi* polypeptide-specific antibodies are used to immunize an animal, preferably a mouse. The splenocytes of such an animal are then used to produce hybridoma cells, and the hybridoma cells are screened to identify clones which produce an antibody whose ability to bind to the *B. burgdorferi* polypeptide-specific antibody can be blocked by the *B. burgdorferi* polypeptide antigen. Such antibodies comprise anti-idiotypic antibodies to the *B. burgdorferi* polypeptide-specific antibody and can be used to immunize an animal to induce formation of further *B. burgdorferi* polypeptide-specific antibodies.

Antibodies and fragements thereof of the present invention may be described by the portion of a polypeptide of the present invention recognized or specifically bound by the antibody. Antibody binding fragements of a polypeptide of the present invention may be described or specified in the same manner as for polypeptide fragements discussed above., i.e, by N-terminal and C-terminal positions or by size in contiguous amino acid residues. Any number of antibody binding fragments, of a polypeptide of the present invention, specified by N-terminal and C-terminal positions or by size in amino acid residues, as described above, may also be excluded from the present invention. Therefore, the present invention includes antibodies the specifically bind a particuarlly discribed fragement of a polypeptide of the present invention and allows for the exclusion of the same.

Antibodies and fragements thereof of the present invention may also be described or specified in terms of their cross-reactivity. Antibodies and fragements that do not bind polypeptides of any other species of *Borrelia* other than *B. burgdorferi* are included in the present invention. Likewise, antibodies and fragements that bind only species of *Borrelia*, i.e. antibodies and fragements that do not bind bacteria from any genus other than *Borrelia*, are included in the present invention.

3. Epitope-Bearing Portions

In another aspect, the invention provides peptides and polypeptides comprising epitope-bearing portions of the *B. burgdorferi* polypeptides of the present invention. These epitopes are immunogenic or antigenic epitopes of the polypeptides of the present invention. An "immunogenic epitope" is defined as a part of a protein that elicits an antibody response when the whole protein or polypeptide is the immunogen. These immunogenic epitopes are believed to be confined to a few loci on the molecule. On the other hand, a region of a protein molecule to which an antibody can bind is defined as an "antigenic determinant" or "antigenic epitope." The number of immunogenic epitopes of a protein generally is less than the number of antigenic epitopes. *See, e.g.,* Geysen, et al. (1983) Proc. Natl. Acad. Sci. USA 81:3998- 4002. Amino acid residues comprising anigenic epitopes may be determined by algorithms such as the the Jameson-Wolf analysis or similar algorithms or by *in vivo* testing for an antigenic response using the methods described herein or those known in the art.

As to the selection of peptides or polypeptides bearing an antigenic epitope (*i.e.*, that contain a region of a protein molecule to which an antibody can bind), it is well known in that art that relatively short synthetic peptides that mimic part of a protein sequence are routinely capable of eliciting an antiserum that reacts with the partially mimicked protein. *See, e.g.,* Sutcliffe, et al., (1983) Science 219:660-666. Peptides capable of eliciting protein-reactive sera are frequently represented in the primary sequence of a protein, can be characterized by a set of simple chemical rules, and are confined neither to immunodominant regions of intact proteins (*i.e.*, immunogenic epitopes) nor to the amino or carboxyl terminals. Peptides that are extremely hydrophobic and those of six or fewer residues generally are ineffective at inducing antibodies that bind to the mimicked protein; longer, peptides, especially those containing proline residues, usually are effective. *See,* Sutcliffe, et al., *supra*, p. 661. For instance, 18 of 20 peptides designed according to these guidelines, containing 8-39 residues covering 75% of the sequence of the influenza virus hemagglutinin HA1 polypeptide chain, induced antibodies that reacted with the HA1 protein or intact virus; and 12/12 peptides from the MuLV polymerase and 18/18 from the rabies glycoprotein induced antibodies that precipitated the respective proteins.

Antigenic epitope-bearing peptides and polypeptides of the invention are therefore useful to raise antibodies, including monoclonal antibodies, that bind specifically to a polypeptide of the invention. Thus, a high proportion of hybridomas obtained by fusion of spleen cells from donors immunized with an antigen epitope-bearing peptide generally secrete antibody reactive with the native protein. *See* Sutcliffe, et al., *supra*, p. 663. The antibodies raised by antigenic epitope-bearing peptides or polypeptides are useful to detect the mimicked protein, and antibodies to different peptides may be used for tracking the fate of various regions of a protein precursor which undergoes post-translational processing. The peptides and anti-peptide antibodies may be used in a variety of qualitative or quantitative assays for the mimicked protein, for instance in competition assays since it has been shown that even short peptides (*e.g.*, about 9 amino acids)

can bind and displace the larger peptides in immunoprecipitation assays. *See, e.g.,* Wilson, et al., (1984) Cell 37:767-778. The anti-peptide antibodies of the invention also are useful for purification of the mimicked protein, for instance, by adsorption chromatography using methods known in the art.

Antigenic epitope-bearing peptides and polypeptides of the invention designed according to the above guidelines preferably contain a sequence of at least seven, more preferably at least nine and most preferably between about 10 to about 50 amino acids (i.e. any integer between 7 and 50) contained within the amino acid sequence of a polypeptide of the invention. However, peptides or polypeptides comprising a larger portion of an amino acid sequence of a polypeptide of the invention, containing about 50 to about 100 amino acids, or any length up to and including the entire amino acid sequence of a polypeptide of the invention, also are considered epitope-bearing peptides or polypeptides of the invention and also are useful for inducing antibodies that react with the mimicked protein. Preferably, the amino acid sequence of the epitope-bearing peptide is selected to provide substantial solubility in aqueous solvents (*i.e.*, the sequence includes relatively hydrophilic residues and highly hydrophobic sequences are preferably avoided); and sequences containing proline residues are particularly preferred.

The epitope-bearing peptides and polypeptides of the present invention may be produced by any conventional means for making peptides or polypeptides including recombinant means using nucleic acid molecules of the invention. For instance, an epitope-bearing amino acid sequence of the present invention may be fused to a larger polypeptide which acts as a carrier during recombinant production and purification, as well as during immunization to produce anti-peptide antibodies. Epitope-bearing peptides also may be synthesized using known methods of chemical synthesis. For instance, Houghten has described a simple method for synthesis of large numbers of peptides, such as 10-20 mg of 248 different 13 residue peptides representing single amino acid variants of a segment of the HA1 polypeptide which were prepared and characterized (by ELISA-type binding studies) in less than four weeks (Houghten, R. A. Proc. Natl. Acad. Sci. USA **82:**5131-5135 (1985)). This "Simultaneous Multiple Peptide Synthesis (SMPS)" process is further described in U.S. Patent No. 4,631,211 to Houghten and coworkers (1986). In this procedure the individual resins for the solid-phase synthesis of various peptides are contained in separate solvent-permeable packets, enabling the optimal use of the many identical repetitive steps involved in solid-phase methods. A completely manual procedure allows 500-1000 or more syntheses to be conducted simultaneously (Houghten et al. (1985) Proc. Natl. Acad. Sci. 82:5131-5135 at 5134.

Epitope-bearing peptides and polypeptides of the invention are used to induce antibodies according to methods well known in the art. *See, e.g.,* Sutcliffe, et al., *supra*;; Wilson, et al., *supra*;; and Bittle, et al. (1985) J. Gen. Virol. 66:2347-2354. Generally, animals may be immunized with free peptide; however, anti-peptide antibody titer may be boosted by coupling of the peptide to a macromolecular carrier, such as keyhole limpet hemacyanin (KLH) or tetanus toxoid. For instance, peptides containing cysteine may be coupled to carrier using a linker such

as m-maleimidobenzoyl-N-hydroxysuccinimide ester (MBS), while other peptides may be coupled to carrier using a more general linking agent such as glutaraldehyde. Animals such as rabbits, rats and mice are immunized with either free or carrier-coupled peptides, for instance, by intraperitoneal and/or intradermal injection of emulsions containing about 100 μg peptide or carrier protein and Freund's adjuvant. Several booster injections may be needed, for instance, at intervals of about two weeks, to provide a useful titer of anti-peptide antibody which can be detected, for example, by ELISA assay using free peptide adsorbed to a solid surface. The titer of anti-peptide antibodies in serum from an immunized animal may be increased by selection of anti-peptide antibodies, for instance, by adsorption to the peptide on a solid support and elution of the selected antibodies according to methods well known in the art.

Immunogenic epitope-bearing peptides of the invention, *i.e.*, those parts of a protein that elicit an antibody response when the whole protein is the immunogen, are identified according to methods known in the art. For instance, Geysen, *et al.*, *supra*, discloses a procedure for rapid concurrent synthesis on solid supports of hundreds of peptides of sufficient purity to react in an ELISA. Interaction of synthesized peptides with antibodies is then easily detected without removing them from the support. In this manner a peptide bearing an immunogenic epitope of a desired protein may be identified routinely by one of ordinary skill in the art. For instance, the immunologically important epitope in the coat protein of foot-and-mouth disease virus was located by Geysen *et al. supra* with a resolution of seven amino acids by synthesis of an overlapping set of all 208 possible hexapeptides covering the entire 213 amino acid sequence of the protein. Then, a complete replacement set of peptides in which all 20 amino acids were substituted in turn at every position within the epitope were synthesized, and the particular amino acids conferring specificity for the reaction with antibody were determined. Thus, peptide analogs of the epitope-bearing peptides of the invention can be made routinely by this method. U.S. Patent No. 4,708,781 to Geysen (1987) further describes this method of identifying a peptide bearing an immunogenic epitope of a desired protein.

Further still, U.S. Patent No. 5,194,392, to Geysen (1990), describes a general method of detecting or determining the sequence of monomers (amino acids or other compounds) which is a topological equivalent of the epitope (*i.e.*, a "mimotope") which is complementary to a particular paratope (antigen binding site) of an antibody of interest. More generally, U.S. Patent No. 4,433,092, also to Geysen (1989), describes a method of detecting or determining a sequence of monomers which is a topographical equivalent of a ligand which is complementary to the ligand binding site of a particular receptor of interest. Similarly, U.S. Patent No. 5,480,971 to Houghten, R. A. *et al.* (1996) discloses linear C_1-C_7-alkyl peralkylated oligopeptides and sets and libraries of such peptides, as well as methods for using such oligopeptide sets and libraries for determining the sequence of a peralkylated oligopeptide that preferentially binds to an acceptor molecule of interest. Thus, non-peptide analogs of the epitope-bearing peptides of the invention also can be made routinely by these methods. The entire disclosure of each document cited in this section on "Polypeptides and Fragments" is

hereby incorporated herein by reference.

As one of skill in the art will appreciate, the polypeptides of the present invention and the epitope-bearing fragments thereof described above can be combined with parts of the constant domain of immunoglobulins (IgG), resulting in chimeric polypeptides. These fusion proteins facilitate purification and show an increased half-life *in vivo*. This has been shown, *e.g.*, for chimeric proteins consisting of the first two domains of the human CD4-polypeptide and various domains of the constant regions of the heavy or light chains of mammalian immunoglobulins. (EPA 0,394,827; Traunecker et al. (1988) Nature 331:84-86. Fusion proteins that have a disulfide-linked dimeric structure due to the IgG part can also be more efficient in binding and neutralizing other molecules than a monomeric *B. burgdorferi* polypeptide or fragment thereof alone. *See* Fountoulakis et al. (1995) J. Biochem. 270:3958-3964. Nucleic acids encoding the above epitopes of *B. burgdorferi* polypeptides can also be recombined with a gene of interest as an epitope tag to aid in detection and purification of the expressed polypeptide.

4. Diagnostic Assays and Kits

The present invention further relates to methods for assaying Borrelia infection in an animal by detecting the expression of genes encoding Borrelia polypeptides of the present invention. The methods comprise analyzing tissue or body fluid from the animal for *Borrelia*-specific antibodies, nucleic acids, or proteins. Analysis of nucleic acid specific to *Borrelia* is assayed by PCR or hybridization techniques using nucleic acid sequences of the present invention as either hybridization probes or primers. *See, e.g.,* Sambrook et al. Molecular cloning: A Laboratory Manual (Cold Spring Harbor Laboratory Press, 2nd ed., 1989, page 54 reference); Eremeeva et al. (1994) J. Clin. Microbiol. 32:803-810 (describing differentiation among spotted fever group *Rickettsiae* species by analysis of restriction fragment length polymorphism of PCR-amplified DNA) and Chen et al. 1994 J. Clin. Microbiol. 32:589-595 (detecting *B burgdorferi* nucleic acids *via* PCR).

Where diagnosis of a disease state related to infection with *Borrelia* has already been made, the present invention is useful for monitoring progression or regression of the disease state whereby patients exhibiting enhanced *Borrelia* gene expression will experience a worse clinical outcome relative to patients expressing these gene(s) at a lower level.

By "biological sample" is intended any biological sample obtained from an animal, cell line, tissue culture, or other source which contains *Borrelia* polypeptide, mRNA, or DNA. Biological samples include body fluids (such as saliva, blood, plasma, urine, mucus, synovial fluid, etc.) tissues (such as muscle, skin, and cartilage) and any other biological source suspected of containing *Borrelia* polypeptides or nucleic acids. Methods for obtaining biological samples such as tissue are well known in the art.

The present invention is useful for detecting diseases related to *Borrelia* infections in animals. Preferred animals include monkeys, apes, cats, dogs, birds, cows, pigs, mice, horses, rabbits and humans. Particularly preferred are humans.

Total RNA can be isolated from a biological sample using any suitable technique such as the single-step guanidinium-thiocyanate-phenol-chloroform method described in Chomczynski et al. (1987) Anal. Biochem. 162:156-159. mRNA encoding *Borrelia* polypeptides having sufficient homology to the nucleic acid sequences identified in SEQ ID NOS:1-155 to allow for hybridization between complementary sequences are then assayed using any appropriate method. These include Northern blot analysis, S1 nuclease mapping, the polymerase chain reaction (PCR), reverse transcription in combination with the polymerase chain reaction (RT-PCR), and reverse transcription in combination with the ligase chain reaction (RT-LCR).

Northern blot analysis can be performed as described in Harada et al. (1990) Cell 63:303-312. Briefly, total RNA is prepared from a biological sample as described above. For the Northern blot, the RNA is denatured in an appropriate buffer (such as glyoxal/dimethyl sulfoxide/sodium phosphate buffer), subjected to agarose gel electrophoresis, and transferred onto a nitrocellulose filter. After the RNAs have been linked to the filter by a UV linker, the filter is prehybridized in a solution containing formamide, SSC, Denhardt's solution, denatured salmon sperm, SDS, and sodium phosphate buffer. A *B. burgdorferi* polynucleotide sequence shown in SEQ ID NOS:1-155 labeled according to any appropriate method (such as the ^{32}P-multiprimed DNA labeling system (Amersham)) is used as probe. After hybridization overnight, the filter is washed and exposed to x-ray film. DNA for use as probe according to the present invention is described in the sections above and will preferably at least 15 nucleotides in length.

S1 mapping can be performed as described in Fujita et al. (1987) Cell 49:357-367. To prepare probe DNA for use in S1 mapping, the sense strand of an above-described *B. burgdorferi* DNA sequence of the present invention is used as a template to synthesize labeled antisense DNA. The antisense DNA can then be digested using an appropriate restriction endonuclease to generate further DNA probes of a desired length. Such antisense probes are useful for visualizing protected bands corresponding to the target mRNA (*i.e.*, mRNA encoding *Borrelia* polypeptides).

Levels of mRNA encoding *Borrelia* polypeptides are assayed, for *e.g.*, using the RT-PCR method described in Makino et al. (1990) Technique 2:295-301. By this method, the radioactivities of the "amplicons" in the polyacrylamide gel bands are linearly related to the initial concentration of the target mRNA. Briefly, this method involves adding total RNA isolated from a biological sample in a reaction mixture containing a RT primer and appropriate buffer. After incubating for primer annealing, the mixture can be supplemented with a RT buffer, dNTPs, DTT, RNase inhibitor and reverse transcriptase. After incubation to achieve reverse transcription of the RNA, the RT products are then subject to PCR using labeled primers. Alternatively, rather than labeling the primers, a labeled dNTP can be included in the PCR reaction mixture. PCR amplification can be performed in a DNA thermal cycler according to conventional techniques. After a suitable number of rounds to achieve amplification, the PCR reaction mixture is electrophoresed on a polyacrylamide gel. After drying the gel, the radioactivity of the appropriate

bands (corresponding to the mRNA encoding the *Borrelia* polypeptides of the present invention) are quantified using an imaging analyzer. RT and PCR reaction ingredients and conditions, reagent and gel concentrations, and labeling methods are well known in the art. Variations on the RT-PCR method will be apparent to the skilled artisan. Other PCR methods that can detect the nucleic acid of the present invention can be found in PCR PRIMER: A LABORATORY MANUAL (C.W. Dieffenbach et al. eds., Cold Spring Harbor Lab Press, 1995).

The polynucleotides of the present invention, including both DNA and RNA, may be used to detect polynucleotides of the present invention or Borrelia species including *B. burgdorferi* using bio chip technology. The present invention includes both high density chip arrays (>1000 oligonucleotides per cm^2) and low density chip arrays (<1000 oligonucleotides per cm^2). Bio chips comprising arrays of polynucleotides of the present invention may be used to detect Borrelia species, including *B. burgdorferi*, in biological and environmental samples and to diagnose an animal, including humans, with an *B. burgdorferi* or other Borrelia infection. The bio chips of the present invention may comprise polynucleotide sequences of other pathogens including bacteria, viral, parasitic, and fungal polynucleotide sequences, in addition to the polynucleotide sequences of the present invention, for use in rapid diffenertial pathogenic detection and diagnosis. The bio chips can also be used to monitor an *B. burgdorferi* or other Borrelia infections and to monitor the genetic changes (deletions, insertions, mismatches, etc.) in response to drug therapy in the clinic and drug development in the laboratory. The bio chip technology comprising arrays of polynucleotides of the present invention may also be used to simultaneously monitor the expression of a multiplicity of genes, including those of the present invention. The polynucleotides used to comprise a selected array may be specified in the same manner as for the fragements, i.e, by their 5' and 3' positions or length in contigious base pairs and include from. Methods and particular uses of the polynucleotides of the present invention to detect Borrelia species, including *B. burgdorferi*, using bio chip technology include those known in the art and those of: U.S. Patent Nos. 5510270, 5545531, 5445934, 5677195, 5532128, 5556752, 5527681, 5451683, 5424186, 5607646, 5658732 and World Patent Nos. WO/9710365, WO/9511995, WO/9743447, WO/9535505, each incorporated herein in their entireties.

Biosensors using the polynucleotides of the present invention may also be used to detect, diagnose, and monitor *B. burgdorferi* or other Borrelia species and infections thereof. Biosensors using the polynucleotides of the present invention may also be used to detect particular polynucleotides of the present invention. Biosensors using the polynucleotides of the present invention may also be used to monitor the genetic changes (deletions, insertions, mismatches, etc.) in response to drug therapy in the clinic and drug development in the laboratory. Methods and particular uses of the polynucleotides of the present invention to detect Borrelia species, including *B. burgdorferi*, using biosenors include those known in the art and those of: U.S. Patent Nos 5721102, 5658732, 5631170, and World Patent Nos. WO97/35011, WO/9720203, each incorporated herein in their entireties.

Thus, the present invention includes both bio chips and biosensors comprising polynucleotides of the present invention and methods of their use.

Assaying *Borrelia* polypeptide levels in a biological sample can occur using any art-known method, such as antibody-based techniques. For example, *Borrelia* polypeptide expression in tissues can be studied with classical immunohistological methods. In these, the specific recognition is provided by the primary antibody (polyclonal or monoclonal) but the secondary detection system can utilize fluorescent, enzyme, or other conjugated secondary antibodies. As a result, an immunohistological staining of tissue section for pathological examination is obtained. Tissues can also be extracted, *e.g.*, with urea and neutral detergent, for the liberation of *Borrelia* polypeptides for Western-blot or dot/slot assay. *See, e.g.,* Jalkanen, M. et al. (1985) J. Cell. Biol. 101:976-985; Jalkanen, M. et al. (1987) J. Cell . Biol. 105:3087-3096. In this technique, which is based on the use of cationic solid phases, quantitation of a *Borrelia* polypeptide can be accomplished using an isolated *Borrelia* polypeptide as a standard. This technique can also be applied to body fluids.

Other antibody-based methods useful for detecting *Borrelia* polypeptide gene expression include immunoassays, such as the ELISA and the radioimmunoassay (RIA). For example, a *Borrelia* polypeptide-specific monoclonal antibodies can be used both as an immunoabsorbent and as an enzyme-labeled probe to detect and quantify a *Borrelia* polypeptide. The amount of a *Borrelia* polypeptide present in the sample can be calculated by reference to the amount present in a standard preparation using a linear regression computer algorithm. Such an ELISA is described in Iacobelli et al. (1988) Breast Cancer Research and Treatment 11:19-30. In another ELISA assay, two distinct specific monoclonal antibodies can be used to detect *Borrelia* polypeptides in a body fluid. In this assay, one of the antibodies is used as the immunoabsorbent and the other as the enzyme-labeled probe.

The above techniques may be conducted essentially as a "one-step" or "two-step" assay. The "one-step" assay involves contacting the *Borrelia* polypeptide with immobilized antibody and, without washing, contacting the mixture with the labeled antibody. The "two-step" assay involves washing before contacting the mixture with the labeled antibody. Other conventional methods may also be employed as suitable. It is usually desirable to immobilize one component of the assay system on a support, thereby allowing other components of the system to be brought into contact with the component and readily removed from the sample. Variations of the above and other immunological methods included in the present invention can also be found in Harlow et al., ANTIBODIES: A LABORATORY MANUAL, (Cold Spring Harbor Laboratory Press, 2nd ed. 1988).

Suitable enzyme labels include, for example, those from the oxidase group, which catalyze the production of hydrogen peroxide by reacting with substrate. Glucose oxidase is particularly preferred as it has good stability and its substrate (glucose) is readily available. Activity of an oxidase label may be assayed by measuring the concentration of hydrogen peroxide formed by the enzyme-labeled antibody/substrate reaction. Besides enzymes, other suitable

labels include radioisotopes, such as iodine (^{125}I, ^{121}I), carbon (^{14}C), sulphur (^{35}S), tritium (^{3}H), indium (^{112}In), and technetium (^{99m}Tc), and fluorescent labels, such as fluorescein and rhodamine, and biotin.

Further suitable labels for the *Borrelia* polypeptide-specific antibodies of the present invention are provided below. Examples of suitable enzyme labels include malate dehydrogenase, Borrelia nuclease, delta-5-steroid isomerase, yeast-alcohol dehydrogenase, alpha-glycerol phosphate dehydrogenase, triose phosphate isomerase, peroxidase, alkaline phosphatase, asparaginase, glucose oxidase, beta-galactosidase, ribonuclease, urease, catalase, glucose-6-phosphate dehydrogenase, glucoamylase, and acetylcholine esterase.

Examples of suitable radioisotopic labels include ^{3}H, ^{111}In, ^{125}I, ^{131}I, ^{32}P, ^{35}S, ^{14}C, ^{51}Cr, ^{57}To, ^{58}Co, ^{59}Fe, ^{75}Se, ^{152}Eu, ^{90}Y, ^{67}Cu, ^{217}Ci, ^{211}At, ^{212}Pb, ^{47}Sc, ^{109}Pd, etc. ^{111}In is a preferred isotope where *in vivo* imaging is used since its avoids the problem of dehalogenation of the ^{125}I or ^{131}I-labeled monoclonal antibody by the liver. In addition, this radionucleotide has a more favorable gamma emission energy for imaging. *See, e.g.*, Perkins et al. (1985) Eur. J. Nucl. Med. 10:296-301; Carasquillo et al. (1987) J. Nucl. Med. 28:281-287. For example, ^{111}In coupled to monoclonal antibodies with 1-(P-isothiocyanatobenzyl)-DPTA has shown little uptake in non-tumors tissues, particularly the liver, and therefore enhances specificity of tumor localization. See, Esteban et al. (1987) J. Nucl. Med. 28:861-870.

Examples of suitable non-radioactive isotopic labels include ^{157}Gd, ^{55}Mn, ^{162}Dy, ^{52}Tr, and ^{56}Fe.

Examples of suitable fluorescent labels include an ^{152}Eu label, a fluorescein label, an isothiocyanate label, a rhodamine label, a phycoerythrin label, a phycocyanin label, an allophycocyanin label, an o-phthaldehyde label, and a fluorescamine label.

Examples of suitable toxin labels include, *Pseudomonas* toxin, diphtheria toxin, ricin, and cholera toxin.

Examples of chemiluminescent labels include a luminal label, an isoluminal label, an aromatic acridinium ester label, an imidazole label, an acridinium salt label, an oxalate ester label, a luciferin label, a luciferase label, and an aequorin label.

Examples of nuclear magnetic resonance contrasting agents include heavy metal nuclei such as Gd, Mn, and iron.

Typical techniques for binding the above-described labels to antibodies are provided by Kennedy et al. (1976) Clin. Chim. Acta 70:1-31, and Schurs et al. (1977) Clin. Chim. Acta 81:1-40. Coupling techniques mentioned in the latter are the glutaraldehyde method, the periodate method, the dimaleimide method, the m-maleimidobenzyl-N-hydroxy-succinimide ester method, all of which methods are incorporated by reference herein.

In a related aspect, the invention includes a diagnostic kit for use in screening serum containing antibodies specific against *B. burgdorferi* infection. Such a kit may include an isolated *B. burgdorferi* antigen comprising an epitope which is specifically immunoreactive with at least one anti-*B. burgdorferi* antibody. Such a kit also includes means for detecting the

binding of said antibody to the antigen. In specific embodiments, the kit may include a recombinantly produced or chemically synthesized peptide or polypeptide antigen. The peptide or polypeptide antigen may be attached to a solid support.

In a more specific embodiment, the detecting means of the above-described kit includes a solid support to which said peptide or polypeptide antigen is attached. Such a kit may also include a non-attached reporter-labeled anti-human antibody. In this embodiment, binding of the antibody to the *B. burgdorferi* antigen can be detected by binding of the reporter labeled antibody to the anti-*B. burgdorferi* polypeptide antibody.

In a related aspect, the invention includes a method of detecting *B. burgdorferi* infection in a subject. This detection method includes reacting a body fluid, preferably serum, from the subject with an isolated *B. burgdorferi* antigen, and examining the antigen for the presence of bound antibody. In a specific embodiment, the method includes a polypeptide antigen attached to a solid support, and serum is reacted with the support. Subsequently, the support is reacted with a reporter-labeled anti-human antibody. The support is then examined for the presence of reporter-labeled antibody.

The solid surface reagent employed in the above assays and kits is prepared by known techniques for attaching protein material to solid support material, such as polymeric beads, dip sticks, 96-well plates or filter material. These attachment methods generally include non-specific adsorption of the protein to the support or covalent attachment of the protein , typically through a free amine group, to a chemically reactive group on the solid support, such as an activated carboxyl, hydroxyl, or aldehyde group. Alternatively, streptavidin coated plates can be used in conjunction with biotinylated antigen(s).

The polypeptides and antibodies of the present invention, including fragments thereof, may be used to detect Borrelia species including *B. burgdorferi* using bio chip and biosensor technology. Bio chip and biosensors of the present invention may comprise the polypeptides of the present invention to detect antibodies, which specifically recognize Borrelia species, including *B. burgdorferi*. Bio chip and biosensors of the present invention may also comprise antibodies which specifically recognize the polypeptides of the present invention to detect Borrelia species, including *B. burgdorferi* or specific polypeptides of the present invention. Bio chips or biosensors comprising polypeptides or antibodies of the present invention may be used to detect Borrelia species, including *B. burgdorferi*, in biological and environmental samples and to diagnose an animal, including humans, with an *B. burgdorferi* or other Borrelia infection. Thus, the present invention includes both bio chips and biosensors comprising polypeptides or antibodies of the present invention and methods of their use.

The bio chips of the present invention may further comprise polypeptide sequences of other pathogens including bacteria, viral, parasitic, and fungal polypeptide sequences, in addition to the polypeptide sequences of the present invention, for use in rapid diffenertial pathogenic detection and diagnosis. The bio chips of the present invention may further comprise antibodies or fragements thereof specific for other pathogens including bacteria, viral, parasitic, and fungal

polypeptide sequences, in addition to the antibodies or fragements thereof of the present invention, for use in rapid diffenertial pathogenic detection and diagnosis. The bio chips and biosensors of the present invention may also be used to monitor an *B. burgdorferi* or other Borrelia infection and to monitor the genetic changes (amio acid deletions, insertions, substitutions, etc.) in response to drug therapy in the clinic and drug development in the laboratory. The bio chip and biosensors comprising polypeptides or antibodies of the present invention may also be used to simultaneously monitor the expression of a multiplicity of polypeptides, including those of the present invention. The polypeptides used to comprise a bio chip or biosensor of the present invention may be specified in the same manner as for the fragements, i.e, by their N-terminal and C-terminal positions or length in contigious amino acid residue. Methods and particular uses of the polypeptides and antibodies of the present invention to detect Borrelia species, including *B. burgdorferi*, or specific polypeptides using bio chip and biosensor technology include those known in the art, those of the U.S. Patent Nos. and World Patent Nos. listed above for bio chips and biosensors using polynucleotides of the present invention, and those of: U.S. Patent Nos. 5658732, 5135852, 5567301, 5677196, 5690894 and World Patent Nos. WO9729366, WO9612957, each incorporated herein in their entireties.

5. Screening Assay for Binding Agents

Using the isolated proteins of the present invention, the present invention further provides methods of obtaining and identifying agents which bind to a protein encoded by one of the ORFs of the present invention or to one of the fragments and the *Borrelia burgdorferi* fragment and contigs herein described.

In general, such methods comprise steps of:

(a) contacting an agent with an isolated protein encoded by one of the ORFs of the present invention, or an isolated fragment of the *Borrelia burgdorferi* genome; and

(b) determining whether the agent binds to said protein or said fragment.

The agents screened in the above assay can be, but are not limited to, peptides, carbohydrates, vitamin derivatives, or other pharmaceutical agents. The agents can be selected and screened at random or rationally selected or designed using protein modeling techniques.

For random screening, agents such as peptides, carbohydrates, pharmaceutical agents and the like are selected at random and are assayed for their ability to bind to the protein encoded by the ORF of the present invention.

Alternatively, agents may be rationally selected or designed. As used herein, an agent is said to be "rationally selected or designed" when the agent is chosen based on the configuration of the particular protein. For example, one skilled in the art can readily adapt currently available procedures to generate peptides, pharmaceutical agents and the like capable of binding to a specific peptide sequence in order to generate rationally designed antipeptide peptides, for example see Hurby *et al.*, "Application of Synthetic Peptides: Antisense Peptides," in *Synthetic*

Peptides, A User's Guide, W. H. Freeman, NY (1992), pp. 289-307, and Kaspczak *et al.*, *Biochemistry 28:*9230-8 (1989), or pharmaceutical agents, or the like.

In addition to the foregoing, one class of agents of the present invention, as broadly described, can be used to control gene expression through binding to one of the ORFs or EMFs of the present invention. As described above, such agents can be randomly screened or rationally designed/selected. Targeting the ORF or EMF allows a skilled artisan to design sequence specific or element specific agents, modulating the expression of either a single ORF or multiple ORFs which rely on the same EMF for expression control.

One class of DNA binding agents are agents which contain base residues which hybridize or form a triple helix by binding to DNA or RNA. Such agents can be based on the classic phosphodiester, ribonucleic acid backbone, or can be a variety of sulfhydryl or polymeric derivatives which have base attachment capacity.

Agents suitable for use in these methods usually contain 20 to 40 bases and are designed to be complementary to a region of the gene involved in transcription (triple helix - see Lee *et al.*, *Nucl. Acids Res. 6:*3073 (1979); Cooney *et al.*, *Science 241:*456 (1988); and Dervan *et al.*, *Science 251:*1360 (1991)) or to the mRNA itself (antisense - Okano, *J. Neurochem. 56:*560 (1991); *Oligodeoxynucleotides as Antisense Inhibitors of Gene Expression,* CRC Press, Boca Raton, FL (1988)). Triple helix- formation optimally results in a shut-off of RNA transcription from DNA, while antisense RNA hybridization blocks translation of an mRNA molecule into polypeptide. Both techniques have been demonstrated to be effective in model systems. Information contained in the sequences of the present invention can be used to design antisense and triple helix-forming oligonucleotides, and other DNA binding agents.

6. Pharmaceutical Compositions and Vaccines

The present invention further provides pharmaceutical agents which can be used to modulate the growth or pathogenicity of *Borrelia burgdorferi*, or another related organism, *in vivo* or *in vitro*. As used herein, a "pharmaceutical agent" is defined as a composition of matter which can be formulated using known techniques to provide a pharmaceutical compositions. As used herein, the "pharmaceutical agents of the present invention" refers the pharmaceutical agents which are derived from the proteins encoded by the ORFs of the present invention or are agents which are identified using the herein described assays.

As used herein, a pharmaceutical agent is said to "modulate the growth pathogenicity of *Borrelia burgdorferi* or a related organism, *in vivo* or *in vitro*," when the agent reduces the rate of growth, rate of division, or viability of the organism in question. The pharmaceutical agents of the present invention can modulate the growth or pathogenicity of an organism in many fashions, although an understanding of the underlying mechanism of action is not needed to practice the use of the pharmaceutical agents of the present invention. Some agents will modulate the growth by binding to an important protein thus blocking the biological activity of the protein, while other agents may bind to a component of the outer surface of the organism blocking attachment or

rendering the organism more prone to act the bodies nature immune system. Alternatively, the agent may comprise a protein encoded by one of the ORFs of the present invention and serve as a vaccine. The development and use of a vaccine based on outer membrane components are well known in the art.

As used herein, a "related organism" is a broad term which refers to any organism whose growth can be modulated by one of the pharmaceutical agents of the present invention. In general, such an organism will contain a homolog of the protein which is the target of the pharmaceutical agent or the protein used as a vaccine. As such, related organisms do not need to be bacterial but may be fungal or viral pathogens.

The pharmaceutical agents and compositions of the present invention may be administered in a convenient manner, such as by the oral, topical, intravenous, intraperitoneal, intramuscular, subcutaneous, intranasal or intradermal routes. The pharmaceutical compositions are administered in an amount which is effective for treating and/or prophylaxis of the specific indication. In general, they are administered in an amount of at least about 1 mg/kg body weight and in most cases they will be administered in an amount not in excess of about 1 g/kg body weight per day. In most cases, the dosage is from about 0.1 mg/kg to about 10 g/kg body weight daily, taking into account the routes of administration, symptoms, *etc.*

The agents of the present invention can be used in native form or can be modified to form a chemical derivative. As used herein, a molecule is said to be a "chemical derivative" of another molecule when it contains additional chemical moieties not normally a part of the molecule. Such moieties may improve the molecule's solubility, absorption, biological half life, *etc.* The moieties may alternatively decrease the toxicity of the molecule, eliminate or attenuate any undesirable side effect of the molecule, *etc.* Moieties capable of mediating such effects are disclosed in, among other sources, REMINGTON'S PHARMACEUTICAL SCIENCES (1980) cited elsewhere herein.

For example, such moieties may change an immunological character of the functional derivative, such as affinity for a given antibody. Such changes in immunomodulation activity are measured by the appropriate assay, such as a competitive type immunoassay. Modifications of such protein properties as redox or thermal stability, biological half-life, hydrophobicity, susceptibility to proteolytic degradation or the tendency to aggregate with carriers or into multimers also may be effected in this way and can be assayed by methods well known to the skilled artisan.

The therapeutic effects of the agents of the present invention may be obtained by providing the agent to a patient by any suitable means (*e.g.*, inhalation, intravenously, intramuscularly, subcutaneously, enterally, or parenterally). It is preferred to administer the agent of the present invention so as to achieve an effective concentration within the blood or tissue in which the growth of the organism is to be controlled. To achieve an effective blood concentration, the preferred method is to administer the agent by injection. The administration may be by continuous infusion, or by single or multiple injections.

In providing a patient with one of the agents of the present invention, the dosage of the administered agent will vary depending upon such factors as the patient's age, weight, height, sex, general medical condition, previous medical history, *etc.* In general, it is desirable to provide the recipient with a dosage of agent which is in the range of from about 1 pg/kg to 10 mg/kg (body weight of patient), although a lower or higher dosage may be administered. The therapeutically effective dose can be lowered by using combinations of the agents of the present invention or another agent.

As used herein, two or more compounds or agents are said to be administered "in combination" with each other when either (1) the physiological effects of each compound, or (2) the serum concentrations of each compound can be measured at the same time. The composition of the present invention can be administered concurrently with, prior to, or following the administration of the other agent.

The agents of the present invention are intended to be provided to recipient subjects in an amount sufficient to decrease the rate of growth (as defined above) of the target organism.

The administration of the agent(s) of the invention may be for either a "prophylactic" or "therapeutic" purpose. When provided prophylactically, the agent(s) are provided in advance of any symptoms indicative of the organisms growth. The prophylactic administration of the agent(s) serves to prevent, attenuate, or decrease the rate of onset of any subsequent infection. When provided therapeutically, the agent(s) are provided at (or shortly after) the onset of an indication of infection. The therapeutic administration of the compound(s) serves to attenuate the pathological symptoms of the infection and to increase the rate of recovery.

The agents of the present invention are administered to a subject, such as a mammal, or a patient, in a pharmaceutically acceptable form and in a therapeutically effective concentration. A composition is said to be "pharmacologically acceptable" if its administration can be tolerated by a recipient patient. Such an agent is said to be administered in a "therapeutically effective amount" if the amount administered is physiologically significant. An agent is physiologically significant if its presence results in a detectable change in the physiology of a recipient patient.

The agents of the present invention can be formulated according to known methods to prepare pharmaceutically useful compositions, whereby these materials, or their functional derivatives, are combined in a mixture with a pharmaceutically acceptable carrier vehicle. Suitable vehicles and their formulation, inclusive of other human proteins, *e.g.*, human serum albumin, are described, for example, in REMINGTON'S PHARMACEUTICAL SCIENCES, 16th Ed., Osol, A., Ed., Mack Publishing, Easton PA (1980). In order to form a pharmaceutically acceptable composition suitable for effective administration, such compositions will contain an effective amount of one or more of the agents of the present invention, together with a suitable amount of carrier vehicle.

Additional pharmaceutical methods may be employed to control the duration of action. Control release preparations may be achieved through the use of polymers to complex or absorb one or more of the agents of the present invention. The controlled delivery may be effectuated by

a variety of well known techniques, including formulation with macromolecules such as, for example, polyesters, polyamino acids, polyvinyl, pyrrolidone, ethylenevinylacetate, methylcellulose, carboxymethylcellulose, or protamine, sulfate, adjusting the concentration of the macromolecules and the agent in the formulation, and by appropriate use of methods of incorporation, which can be manipulated to effectuate a desired time course of release. Another possible method to control the duration of action by controlled release preparations is to incorporate agents of the present invention into particles of a polymeric material such as polyesters, polyamino acids, hydrogels, poly(lactic acid) or ethylene vinylacetate copolymers. Alternatively, instead of incorporating these agents into polymeric particles, it is possible to entrap these materials in microcapsules prepared, for example, by coacervation techniques or by interfacial polymerization with, for example, hydroxymethylcellulose or gelatine-microcapsules and poly(methylmethacylate) microcapsules, respectively, or in colloidal drug delivery systems, for example, liposomes, albumin microspheres, microemulsions, nanoparticles, and nanocapsules or in macroemulsions. Such techniques are disclosed in REMINGTON'S PHARMACEUTICAL SCIENCES (1980).

The invention further provides a pharmaceutical pack or kit comprising one or more containers filled with one or more of the ingredients of the pharmaceutical compositions of the invention. Associated with such container(s) can be a notice in the form prescribed by a governmental agency regulating the manufacture, use or sale of pharmaceuticals or biological products, which notice reflects approval by the agency of manufacture, use or sale for human administration.

In addition, the agents of the present invention may be employed in conjunction with other therapeutic compounds.

7. Shot-Gun Approach to Megabase DNA Sequencing

The present invention further demonstrates that a large sequence can be sequenced using a random shotgun approach. This procedure, described in detail in the examples that follow, has eliminated the up front cost of isolating and ordering overlapping or contiguous subclones prior to the start of the sequencing protocols.

Certain aspects of the present invention are described in greater detail in the examples that follow. The examples are provided by way of illustration. Other aspects and embodiments of the present invention are contemplated by the inventors, as will be clear to those of skill in the art from reading the present disclosure.

ILLUSTRATIVE EXAMPLES

LIBRARIES AND SEQUENCING

1. Shotgun Sequencing Probability Analysis

The overall strategy for a shotgun approach to whole genome sequencing follows from the Lander and Waterman (Landerman and Waterman, *Genomics* 2:231 (1988)) application of the equation for the Poisson distribution. According to this treatment, the probability, P0, that any given base in a sequence of size L, in nucleotides, is not sequenced after a certain amount, n, in nucleotides, of random sequence has been determined can be calculated by the equation P0 = e-m, where m is L/n, the fold coverage. For instance, for a genome of 2.8 Mb, m=1 when 2.8 Mb of sequence has been randomly generated (1X coverage). At that point, P0 = e-1 = 0.37. The probability that any given base has not been sequenced is the same as the probability that any region of the whole sequence L has not been determined and, therefore, is equivalent to the fraction of the whole sequence that has yet to be determined. Thus, at one-fold coverage, approximately 37% of a polynucleotide of size L, in nucleotides has not been sequenced. When 14 Mb of sequence has been generated, coverage is 5X for a 2.8 Mb and the unsequenced fraction drops to .0067 or 0.67%. 5X coverage of a 2.8 Mb sequence can be attained by sequencing approximately 17,000 random clones from both insert ends with an average sequence read length of 410 bp.

Similarly, the total gap length, G, is determined by the equation G = Le-m, and the average gap size, g, follows the equation, g = L/n. Thus, 5X coverage leaves about 240 gaps averaging about 82 bp in size in a sequence of a polynucleotide 2.8 Mb long.

The treatment above is essentially that of Lander and Waterman, *Genomics* 2: 231 (1988).

2. Random Library Construction

In order to approximate the random model described above during actual sequencing, a nearly ideal library of cloned genomic fragments is required. The following library construction procedure was developed to achieve this end.

Borrelia burgdorferi DNA is prepared by phenol extraction. A mixture containing 200 µg DNA in 1.0 ml of 300 mM sodium acetate, 10 mM Tris-HCl, 1 mM Na-EDTA, 50% glycerol is processed through a nebulizer (IPI Medical Products) with a stream of nitrogen adjusted to 35 Kpa for 2 minutes. The sonicated DNA is ethanol precipitated and redissolved in 500 µl TE buffer.

To create blunt-ends, a 100 µl aliquot of the resuspended DNA is digested with 5 units of BAL31 nuclease (New England BioLabs) for 10 min at 30°C in 200 µl BAL31 buffer. The digested DNA is phenol-extracted, ethanol-precipitated, redissolved in 100 µl TE buffer, and then size-fractionated by electrophoresis through a 1.0% low melting temperature agarose gel. The section containing DNA fragments 1.6-2.0 kb in size is excised from the gel, and the LGT agarose is melted and the resulting solution is extracted with phenol to separate the agarose from the DNA. DNA is ethanol precipitated and redissolved in 20 µl of TE buffer for ligation to vector.

A two-step ligation procedure is used to produce a plasmid library with 97% inserts, of which >99% were single inserts. The first ligation mixture (50 ul) contains 2 µg of DNA fragments, 2 µg pUC18 DNA (Pharmacia) cut with SmaI and dephosphorylated with bacterial alkaline phosphatase, and 10 units of T4 ligase (GIBCO/BRL) and is incubated at 14°C for 4 hr. The ligation mixture then is phenol extracted and ethanol precipitated, and the precipitated DNA is dissolved in 20 µl TE buffer and electrophoresed on a 1.0% low melting agarose gel. Discrete bands in a ladder are visualized by ethidium bromide-staining and UV illumination and identified by size as insert (I), vector (v), v+I, v+2i, v+3i, *etc.* The portion of the gel containing v+I DNA is excised and the v+I DNA is recovered and resuspended into 20 µl TE. The v+I DNA then is blunt-ended by T4 polymerase treatment for 5 min. at 37°C in a reaction mixture (50 ul) containing the v+I linears, 500 µM each of the 4 dNTPs, and 9 units of T4 polymerase (New England BioLabs), under recommended buffer conditions. After phenol extraction and ethanol precipitation the repaired v+I linears are dissolved in 20 µl TE. The final ligation to produce circles is carried out in a 50 µl reaction containing 5 µl of v+I linears and 5 units of T4 ligase at 14°C overnight. After 10 min. at 70°C the following day, the reaction mixture is stored at -20°C.

This two-stage procedure results in a molecularly random collection of single-insert plasmid recombinants with minimal contamination from double-insert chimeras (<1%) or free vector (<3%).

Since deviation from randomness can arise from propagation the DNA in the host, *E. coli* host cells deficient in all recombination and restriction functions (A. Greener, *Strategies 3 (1)*:5 (1990)) are used to prevent rearrangements, deletions, and loss of clones by restriction. Furthermore, transformed cells are plated directly on antibiotic diffusion plates to avoid the usual broth recovery phase which allows multiplication and selection of the most rapidly growing cells.

Plating is carried out as follows. A 100 µl aliquot of Epicurian Coli SURE II Supercompetent Cells (Stratagene 200152) is thawed on ice and transferred to a chilled Falcon 2059 tube on ice. A 1.7 µl aliquot of 1.42 M beta-mercaptoethanol is added to the aliquot of cells to a final concentration of 25 mM. Cells are incubated on ice for 10 min. A 1 µl aliquot of the final ligation is added to the cells and incubated on ice for 30 min. The cells are heat pulsed for 30 sec. at 42°C and placed back on ice for 2 min. The outgrowth period in liquid culture is eliminated from this protocol in order to minimize the preferential growth of any given transformed cell. Instead the transformation mixture is plated directly on a nutrient rich SOB plate containing a 5 ml bottom layer of SOB agar (5% SOB agar: 20 g tryptone, 5 g yeast extract, 0.5 g NaCl, 1.5% Difco Agar per liter of media). The 5 ml bottom layer is supplemented with 0.4 ml of 50 mg/ml ampicillin per 100 ml SOB agar. The 15 ml top layer of SOB agar is supplemented with 1 ml X-Gal (2%), 1 ml MgCl2 (1 M), and 1 ml MgSO4/100 ml SOB agar. The 15 ml top layer is poured just prior to plating. Our titer is approximately 100 colonies/10 µl aliquot of transformation.

All colonies are picked for template preparation regardless of size. Thus, only clones lost due to "poison" DNA or deleterious gene products are deleted from the library, resulting in a slight increase in gap number over that expected.

3. Random DNA Sequencing

High quality double stranded DNA plasmid templates are prepared using a "boiling bead" method developed in collaboration with Advanced Genetic Technology Corp. (Gaithersburg, MD) (Adams *et al.*, *Science 252*:1651 (1991); Adams *et al.*, *Nature 355*:632 (1992)). Plasmid preparation is performed in a 96-well format for all stages of DNA preparation from bacterial growth through final DNA purification. Template concentration is determined using Hoechst Dye and a Millipore Cytofluor. DNA concentrations are not adjusted, but low-yielding templates are identified where possible and not sequenced.

Templates are also prepared from two *Borrelia burgdorferi* lambda genomic libraries. An amplified library is constructed in the vector Lambda GEM-12 (Promega) and an unamplified library is constructed in Lambda DASH II (Stratagene). In particular, for the unamplified lambda library, *Borrelia burgdorferi* DNA (> 100 kb) is partially digested in a reaction mixture (200 ul) containing 50 µg DNA, 1X Sau3AI buffer, 20 units Sau3AI for 6 min. at 23°C. The digested DNA was phenol-extracted and electrophoresed on a 0.5% low melting agarose gel at 2V/cm for 7 hours. Fragments from 15 to 25 kb are excised and recovered in a final volume of 6 ul. One µl of fragments is used with 1 µl of DASHII vector (Stratagene) in the recommended ligation reaction. One µl of the ligation mixture is used per packaging reaction following the recommended protocol with the Gigapack II XL Packaging Extract (Stratagene, #227711). Phage are plated directly without amplification from the packaging mixture (after dilution with 500 µl of recommended SM buffer and chloroform treatment). Yield is about 2.5x103 pfu/ul. The amplified library is prepared essentially as above except the lambda GEM-12 vector is used. After packaging, about 3.5x104 pfu are plated on the restrictive NM539 host. The lysate is harvested in 2 ml of SM buffer and stored frozen in 7% dimethylsulfoxide. The phage titer is approximately 1x109 pfu/ml.

Liquid lysates (100 µl) are prepared from randomly selected plaques (from the unamplified library) and template is prepared by long-range PCR using T7 and T3 vector-specific primers.

Sequencing reactions are carried out on plasmid and/or PCR templates using the AB Catalyst LabStation with Applied Biosystems PRISM Ready Reaction Dye Primer Cycle Sequencing Kits for the M13 forward (M13-21) and the M13 reverse (M13RP1) primers (Adams *et al.*, *Nature 368*:474 (1994)). Dye terminator sequencing reactions are carried out on the lambda templates on a Perkin-Elmer 9600 Thermocycler using the Applied Biosystems Ready Reaction Dye Terminator Cycle Sequencing kits. T7 and SP6 primers are used to sequence the ends of the inserts from the Lambda GEM-12 library and T7 and T3 primers are used to sequence the ends of the inserts from the Lambda DASH II library. Sequencing reactions are performed

by eight individuals using an average of fourteen AB 373 DNA Sequencers per day. All sequencing reactions are analyzed using the Stretch modification of the AB 373, primarily using a 34 cm well-to-read distance. The overall sequencing success rate very approximately is about 85% for M13-21 and M13RP1 sequences and 65% for dye-terminator reactions. The average usable read length is 485 bp for M13-21 sequences, 445bp for M13RP1 sequences, and 375 bp for dye-terminator reactions.

Richards *et al.,* Chapter 28 in AUTOMATED DNA SEQUENCING AND ANALYSIS, M. D. Adams, C. Fields, J. C. Venter, Eds., Academic Press, London, (1994) described the value of using sequence from both ends of sequencing templates to facilitate ordering of contigs in shotgun assembly projects of lambda and cosmid clones. We balance the desirability of both-end sequencing (including the reduced cost of lower total number of templates) against shorter read-lengths for sequencing reactions performed with the M13RP1 (reverse) primer compared to the M13-21 (forward) primer. Approximately one-half of the templates are sequenced from both ends. Random reverse sequencing reactions are done based on successful forward sequencing reactions. Some M13RP1 sequences are obtained in a semi-directed fashion: M13-21: sequences pointing outward at the ends of contigs are chosen for M13RP1 sequencing in an effort to specifically order contigs.

4. Protocol for Automated Cycle Sequencing

The sequencing is carried out using ABI Catalyst robots and AB 373 Automated DNA Sequencers. The Catalyst robot is a publicly available sophisticated pipetting and temperature control robot which has been developed specifically for DNA sequencing reactions. The Catalyst combines pre-aliquoted templates and reaction mixes consisting of deoxy- and dideoxynucleotides, the thermostable Taq DNA polymerase, fluorescently-labelled sequencing primers, and reaction buffer. Reaction mixes and templates are combined in the wells of an aluminum 96-well thermocycling plate. Thirty consecutive cycles of linear amplification (i.*e.*., one primer synthesis) steps are performed including denaturation, annealing of primer and template, and extension; *i.e.*, DNA synthesis. A heated lid with rubber gaskets on the thermocycling plate prevents evaporation without the need for an oil overlay.

Two sequencing protocols are used: one for dye-labelled primers and a second for dye-labelled dideoxy chain terminators. The shotgun sequencing involves use of four dye-labelled sequencing primers, one for each of the four terminator nucleotide. Each dye-primer is labelled with a different fluorescent dye, permitting the four individual reactions to be combined into one lane of the 373 DNA Sequencer for electrophoresis, detection, and base-calling. ABI currently supplies pre-mixed reaction mixes in bulk packages containing all the necessary non-template reagents for sequencing. Sequencing can be done with both plasmid and PCR- generated templates with both dye-primers and dye- terminators with approximately equal fidelity, although plasmid templates generally give longer usable sequences.

Thirty-two reactions are loaded per AB373 Sequencer each day, for a total of 960 samples. Electrophoresis is run overnight following the manufacturer's protocols, and the data is collected for twelve hours. Following electrophoresis and fluorescence detection, the ABI 373 performs automatic lane tracking and base-calling. The lane-tracking is confirmed visually. Each sequence electropherogram (or fluorescence lane trace) is inspected visually and assessed for quality. Trailing sequences of low quality are removed and the sequence itself is loaded via software to a Sybase database (archived daily to 8mm tape). Leading vector polylinker sequence is removed automatically by a software program. Average edited lengths of sequences from the standard ABI 373 are around 400 bp and depend mostly on the quality of the template used for the sequencing reaction. ABI 373 Sequencers converted to Stretch Liners provide a longer electrophoresis path prior to fluorescence detection and increase the average number of usable bases to 500-600 bp.

INFORMATICS

1. Data Management

A number of information management systems for a large-scale sequencing lab have been developed. (For review see, for instance, Kerlavage *et al.*, *Proceedings of the Twenty-Sixth Annual Hawaii International Conference on System Sciences*, IEEE Computer Society Press, Washington D. C., 585 (1993)) The system used to collect and assemble the sequence data was developed using the Sybase relational database management system and was designed to automate data flow wherever possible and to reduce user error. The database stores and correlates all information collected during the entire operation from template preparation to final analysis of the genome. Because the raw output of the ABI 373 Sequencers was based on a Macintosh platform and the data management system chosen was based on a Unix platform, it was necessary to design and implement a variety of multi- user, client-server applications which allow the raw data as well as analysis results to flow seamlessly into the database with a minimum of user effort.

2. Assembly

An assembly engine (TIGR Assembler) developed for the rapid and accurate assembly of thousands of sequence fragments was employed to generate contigs. The TIGR assembler simultaneously clusters and assembles fragments of the genome. In order to obtain the speed necessary to assemble more than 104 fragments, the algorithm builds a hash table of 12 bp oligonucleotide subsequences to generate a list of potential sequence fragment overlaps. The number of potential overlaps for each fragment determines which fragments are likely to fall into repetitive elements. Beginning with a single seed sequence fragment, TIGR Assembler extends the current contig by attempting to add the best matching fragment based on oligonucleotide content. The contig and candidate fragment are aligned using a modified version of the Smith-Waterman algorithm which provides for optimal gapped alignments (Waterman, M. S., *Methods*

*in Enzymology 164:*765 (1988)). The contig is extended by the fragment only if strict criteria for the quality of the match are met. The match criteria include the minimum length of overlap, the maximum length of an unmatched end, and the minimum percentage match. These criteria are automatically lowered by the algorithm in regions of minimal coverage and raised in regions with a possible repetitive element. The number of potential overlaps for each fragment determines which fragments are likely to fall into repetitive elements. Fragments representing the boundaries of repetitive elements and potentially chimeric fragments are often rejected based on partial mismatches at the ends of alignments and excluded from the current contig. TIGR Assembler is designed to take advantage of clone size information coupled with sequencing from both ends of each template. It enforces the constraint that sequence fragments from two ends of the same template point toward one another in the contig and are located within a certain range of base pairs (definable for each clone based on the known clone size range for a given library). The process resulted in 155 contigs as represented by SEQ ID NOs:1-155.

3. Identifying Genes

The predicted coding regions of the *Borrelia burgdorferi* genome were initially defined with the program GeneMark, which finds ORFs using a probabilistic classification technique. The predicted coding region sequences were used in searches against a database of all nucleotide sequences from GenBank (July, 1997), using the BLASTN search method to identify overlaps of 50 or more nucleotides with at least a 95% identity (using default parameters). Those ORFs with nucleotide sequence matches are shown in Table 1. The ORFs without such matches were translated to protein sequences and compared to a non-redundant database of known proteins generated by combining the Swiss-prot, PIR and GenPept databases. ORFs that matched a database protein with BLASTP probability less than or equal to 0.01 are shown in Table 2. The table also lists assigned functions based on the closest match in the databases. ORFs that did not match protein or nucleotide sequences in the databases at these levels are shown in Table 3.

ILLUSTRATIVE APPLICATIONS

1. Production of an Antibody to a *Borrelia burgdorferi* Protein

Substantially pure protein or polypeptide is isolated from the transfected or transformed cells using any one of the methods known in the art. The protein can also be produced in a recombinant prokaryotic expression system, such as *E. coli*, or can be chemically synthesized. Concentration of protein in the final preparation is adjusted, for example, by concentration on an Amicon filter device, to the level of a few micrograms/ml. Monoclonal or polyclonal antibody to the protein can then be prepared as follows.

2. Monoclonal Antibody Production by Hybridoma Fusion

Monoclonal antibody to epitopes of any of the peptides identified and isolated as described can be prepared from murine hybridomas according to the classical method of Kohler, G. and Milstein, C., *Nature 256*:495 (1975) or modifications of the methods thereof. Briefly, a mouse is repetitively inoculated with a few micrograms of the selected protein over a period of a few weeks. The mouse is then sacrificed, and the antibody producing cells of the spleen isolated. The spleen cells are fused by means of polyethylene glycol with mouse myeloma cells, and the excess unfused cells destroyed by growth of the system on selective media comprising aminopterin (HAT media). The successfully fused cells are diluted and aliquots of the dilution placed in wells of a microtiter plate where growth of the culture is continued. Antibody-producing clones are identified by detection of antibody in the supernatant fluid of the wells by immunoassay procedures, such as ELISA, as originally described by Engvall, E., *Meth. Enzymol. 70:*419 (1980), and modified methods thereof. Selected positive clones can be expanded and their monoclonal antibody product harvested for use. Detailed procedures for monoclonal antibody production are described in Davis, L. *et al.*, *Basic Methods in Molecular Biology,* Elsevier, New York. Section 21-2 (1989).

3. Polyclonal Antibody Production by Immunization

Polyclonal antiserum containing antibodies to heterogenous epitopes of a single protein can be prepared by immunizing suitable animals with the expressed protein described above, which can be unmodified or modified to enhance immunogenicity. Effective polyclonal antibody production is affected by many factors related both to the antigen and the host species. For example, small molecules tend to be less immunogenic than others and may require the use of carriers and adjuvant. Also, host animals vary in response to site of inoculations and dose, with both inadequate or excessive doses of antigen resulting in low titer antisera. Small doses (ng level) of antigen administered at multiple intradermal sites appears to be most reliable. An effective immunization protocol for rabbits can be found in Vaitukaitis, J. *et al.*, *J. Clin.* Endocrinol. Metab. 33:988-991 (1971).

Booster injections can be given at regular intervals, and antiserum harvested when antibody titer thereof, as determined semi-quantitatively, for example, by double immunodiffusion in agar against known concentrations of the antigen, begins to fall. See, for example, Ouchterlony, O. *et al.*, Chap. 19 in: *Handbook of Experimental Immunology,* Wier, D., ed, Blackwell (1973). Plateau concentration of antibody is usually in the range of 0. 1 to 0. 2 mg/ml of serum (about 12M). Affinity of the antisera for the antigen is determined by preparing competitive binding curves, as described, for example, by Fisher, D., Chap. 42 in: *Manual of Clinical Immunology*, second edition, Rose and Friedman, eds., Amer. Soc. For Microbiology, Washington, D. C. (1980)

Antibody preparations prepared according to either protocol are useful in quantitative immunoassays which determine concentrations of antigen-bearing substances in biological

samples; they are also used semi- quantitatively or qualitatively to identify the presence of antigen in a biological sample. In addition, antibodies are useful in various animal models of pneumococcal disease as a means of evaluating the protein used to make the antibody as a potential vaccine target or as a means of evaluating the antibody as a potential immunotherapeutic or immunoprophylactic reagent.

4. Preparation of PCR Primers and Amplification of DNA

Various fragments of the *Borrelia burgdorferi* genome, such as those of Tables 1-6 and SEQ ID NOS: 1-155 can be used, in accordance with the present invention, to prepare PCR primers for a variety of uses. The PCR primers are preferably at least 15 bases, and more preferably at least 18 bases in length. When selecting a primer sequence, it is preferred that the primer pairs have approximately the same G/C ratio, so that melting temperatures are approximately the same. The PCR primers and amplified DNA of this Example find use in the Examples that follow.

5. Isolation of a Selected DNA Clone From B. burgdorferi

Three approaches are used to isolate a *B. burgdorferi* clone comprising a polynucleotide of the present invention from any *B. burgdorferi* genomic DNA library. The *B. burgdorferi* strain B31PU has been deposited as a convienent source for obtaining a *B. burgdorferi* strain although a wide varity of strains *B. burgdorferi* strains can be used which are known in the art.

B. burgdorferi genomic DNA is prepared using the following method. A 20ml overnight bacterial culture grown in a rich medium (e.g., Trypticase Soy Broth, Brain Heart Infusion broth or Super broth), pelleted, ished two times with TES (30mM Tris-pH 8.0, 25mM EDTA, 50mM NaCl), and resuspended in 5ml high salt TES (2.5M NaCl). Lysostaphin is added to final concentration of approx 50ug/ml and the mixture is rotated slowly 1 hour at 37C to make protoplast cells. The solution is then placed in incubator (or place in a shaking water bath) and warmed to 55C. Five hundred micro liter of 20% sarcosyl in TES (final concentration 2%) is then added to lyse the cells. Next, guanidine HCl is added to a final concentration of 7M (3.69g in 5.5 ml). The mixture is swirled slowly at 55C for 60-90 min (solution should clear). A CsCl gradient is then set up in SW41 ultra clear tubes using 2.0ml 5.7M CsCl and overlaying with 2.85M CsCl. The gradient is carefully overlayed with the DNA-containing GuHCl solution. The gradient is spun at 30,000 rpm, 20C for 24 hr and the lower DNA band is collected. The volume is increased to 5 ml with TE buffer. The DNA is then treated with protease K (10 ug/ml) overnight at 37 C, and precipitated with ethanol. The precipitated DNA is resuspended in a desired buffer.

In the first method, a plasmid is directly isolated by screening a plasmid *B. burgdorferi* genomic DNA library using a polynucleotide probe corresponding to a polynucleotide of the present invention. Particularly, a specific polynucleotide with 30-40 nucleotides is synthesized using an Applied Biosystems DNA synthesizer according to the sequence reported. The

oligonucleotide is labeled, for instance, with ^{32}P-γ-ATP using T4 polynucleotide kinase and purified according to routine methods. (*See, e.g.*, Maniatis et al., Molecular Cloning: A Laboratory Manual, Cold Spring Harbor Press, Cold Spring, NY (1982).) The library is transformed into a suitable host, as indicated above (such as XL-1 Blue (Stratagene)) using techniques known to those of skill in the art. *See, e.g.*, Sambrook et al. MOLECULAR CLONING: A LABORATORY MANUAL (Cold Spring Harbor, N.Y. 2nd ed. 1989); Ausubel et al., CURRENT PROTOCALS IN MOLECULAR BIOLOGY (John Wiley and Sons, N.Y. 1989). The transformants are plated on 1.5% agar plates (containing the appropriate selection agent, e.g., ampicillin) to a density of about 150 transformants (colonies) per plate. These plates are screened using Nylon membranes according to routine methods for bacterial colony screening. *See, e.g.*, Sambrook et al. MOLECULAR CLONING: A LABORATORY MANUAL (Cold Spring Harbor, N.Y. 2nd ed. 1989); Ausubel et al., CURRENT PROTOCALS IN MOLECULAR BIOLOGY (John Wiley and Sons, N.Y. 1989) or other techniques known to those of skill in the art.

Alternatively, two primers of 15-25 nucleotides derived from the 5' and 3' ends of a polynucleotide of SEQ ID NOS:1-155 are synthesized and used to amplify the desired DNA by PCR using a *B. burgdorferi* genomic DNA prep as a template. PCR is carried out under routine conditions, for instance, in 25 µl of reaction mixture with 0.5 ug of the above DNA template. A convenient reaction mixture is 1.5-5 mM $MgCl_2$, 0.01% (w/v) gelatin, 20 µM each of dATP, dCTP, dGTP, dTTP, 25 pmol of each primer and 0.25 Unit of Taq polymerase. Thirty five cycles of PCR (denaturation at 94°C for 1 min; annealing at 55°C for 1 min; elongation at 72°C for 1 min) are performed with a Perkin-Elmer Cetus automated thermal cycler. The amplified product is analyzed by agarose gel electrophoresis and the DNA band with expected molecular weight is excised and purified. The PCR product is verified to be the selected sequence by subcloning and sequencing the DNA product.

Finally, overlapping oligos of the DNA sequences of SEQ ID NOS:1-155 can be chemically synthesized and used to generate a nucleotide sequence of desired length using PCR methods known in the art.

6(a). Expression and Purification Borrelia polypeptides in E. coli

The bacterial expression vector pQE60 is used for bacterial expression of some of the polypeptide fragements of the present invention. (QIAGEN, Inc., 9259 Eton Avenue, Chatsworth, CA, 91311). pQE60 encodes ampicillin antibiotic resistance ("Ampr") and contains a bacterial origin of replication ("ori"), an IPTG inducible promoter, a ribosome binding site ("RBS"), six codons encoding histidine residues that allow affinity purification using nickel-nitrilo-tri-acetic acid ("Ni-NTA") affinity resin (QIAGEN, Inc., *supra*) and suitable single restriction enzyme cleavage sites. These elements are arranged such that an inserted DNA fragment encoding a polypeptide expresses that polypeptide with the six His residues (i.e., a "6

X His tag") covalently linked to the carboxyl terminus of that polypeptide.

The DNA sequence encoding the desired portion of a *B. burgdorferi* protein of the present invention is amplified from *B. burgdorferi* genomic DNA using PCR oligonucleotide primers which anneal to the 5' and 3' sequences coding for the portions of the *B. burgdorferi* polynucleotide shown in SEQ ID NOS:1-155. Additional nucleotides containing restriction sites to facilitate cloning in the pQE60 vector are added to the 5' and 3' sequences, respectively.

For cloning the mature protein, the 5' primer has a sequence containing an appropriate restriction site followed by nucleotides of the amino terminal coding sequence of the desired *B. burgdorferi* polynucleotide sequence in SEQ ID NOS:1-155. One of ordinary skill in the art would appreciate that the point in the protein coding sequence where the 5' and 3' primers begin may be varied to amplify a DNA segment encoding any desired portion of the complete protein shorter or longer than the mature form. The 3' primer has a sequence containing an appropriate restriction site followed by nucleotides complementary to the 3' end of the polypeptide coding sequence of SEQ ID NOS:1-155, excluding a stop codon, with the coding sequence aligned with the restriction site so as to maintain its reading frame with that of the six His codons in the pQE60 vector.

The amplified *B. burgdorferi* DNA fragment and the vector pQE60 are digested with restriction enzymes which recognize the sites in the primers and the digested DNAs are then ligated together. The *B. burgdorferi* DNA is inserted into the restricted pQE60 vector in a manner which places the *B. burgdorferi* protein coding region downstream from the IPTG-inducible promoter and in-frame with an initiating AUG and the six histidine codons.

The ligation mixture is transformed into competent *E. coli* cells using standard procedures such as those described by Sambrook et al., *supra*.. *E. coli* strain M15/rep4, containing multiple copies of the plasmid pREP4, which expresses the lac repressor and confers kanamycin resistance ("Kanr"), is used in carrying out the illustrative example described herein. This strain, which is only one of many that are suitable for expressing a *B. burgdorferi* polypeptide, is available commercially (QIAGEN, Inc., *supra*). Transformants are identified by their ability to grow on LB agar plates in the presence of ampicillin and kanamycin. Plasmid DNA is isolated from resistant colonies and the identity of the cloned DNA confirmed by restriction analysis, PCR and DNA sequencing.

Clones containing the desired constructs are grown overnight ("O/N") in liquid culture in LB media supplemented with both ampicillin (100 µg/ml) and kanamycin (25 µg/ml). The O/N culture is used to inoculate a large culture, at a dilution of approximately 1:25 to 1:250. The cells are grown to an optical density at 600 nm ("OD600") of between 0.4 and 0 6. Isopropyl-β-D-thiogalactopyranoside ("IPTG") is then added to a final concentration of 1 mM to induce transcription from the lac repressor sensitive promoter, by inactivating the lacI repressor. Cells subsequently are incubated further for 3 to 4 hours. Cells then are harvested by centrifugation.

The cells are then stirred for 3-4 hours at 4°C in 6M guanidine-HCl, pH 8. The cell

debris is removed by centrifugation, and the supernatant containing the *B. burgdorferi* polypeptide is loaded onto a nickel-nitrilo-tri-acetic acid ("Ni-NTA") affinity resin column (QIAGEN, Inc., *supra*). Proteins with a 6 x His tag bind to the Ni-NTA resin with high affinity are purified in a simple one-step procedure (for details see: The QIAexpressionist, 1995, QIAGEN, Inc., *supra*). Briefly the supernatant is loaded onto the column in 6 M guanidine-HCl, pH 8, the column is first washed with 10 volumes of 6 M guanidine-HCl, pH 8, then washed with 10 volumes of 6 M guanidine-HCl pH 6, and finally the *B. burgdorferi* polypeptide is eluted with 6 M guanidine-HCl, pH 5.

The purified protein is then renatured by dialyzing it against phosphate-buffered saline (PBS) or 50 mM Na-acetate, pH 6 buffer plus 200 mM NaCl. Alternatively, the protein could be successfully refolded while immobilized on the Ni-NTA column. The recommended conditions are as follows: renature using a linear 6M-1M urea gradient in 500 mM NaCl, 20% glycerol, 20 mM Tris/HCl pH 7.4, containing protease inhibitors. The renaturation should be performed over a period of 1.5 hours or more. After renaturation the proteins can be eluted by the addition of 250 mM immidazole. Immidazole is removed by a final dialyzing step against PBS or 50 mM sodium acetate pH 6 buffer plus 200 mM NaCl. The purified protein is stored at 4° C or frozen at -80° C.

The polypeptide of the present invention are also prepared using a non-denaturing protein purification method. For these polypeptides, the cell pellet from each liter of culture is resuspended in 25 mls of Lysis Buffer A at 4°C (Lysis Buffer A = 50 mM Na-phosphate, 300 mM NaCl, 10 mM 2-mercaptoethanol, 10% Glycerol, pH 7.5 with 1 tablet of Complete EDTA-free protease inhibitor cocktail (Boehringer Mannheim #1873580) per 50 ml of buffer). Absorbance at 550 nm is approximately 10-20 O.D./ml. The suspension is then put through three freeze/thaw cycles from -70°C (using a ethanol-dry ice bath) up to room temperature. The cells are lysed via sonication in short 10 sec bursts over 3 minutes at approximately 80W while kept on ice. The sonicated sample is then centrifuged at 15,000 RPM for 30 minutes at 4°C. The supernatant is passed through a column containing 1.0 ml of CL-4B resin to pre-clear the sample of any proteins that may bind to agarose non-specifically, and the flow-through fraction is collected.

The pre-cleared flow-through is applied to a nickel-nitrilo-tri-acetic acid ("Ni-NTA") affinity resin column (Quiagen, Inc., *supra*). Proteins with a 6 X His tag bind to the Ni-NTA resin with high affinity and can be purified in a simple one-step procedure. Briefly, the supernatant is loaded onto the column in Lysis Buffer A at 4°C, the column is first washed with 10 volumes of Lysis Buffer A until the A280 of the eluate returns to the baseline. Then, the column is washed with 5 volumes of 40 mM Imidazole (92% Lysis Buffer A / 8% Buffer B) (Buffer B = 50 mM Na-Phosphate, 300 mM NaCl, 10% Glycerol, 10 mM 2-mercaptoethanol, 500 mM Imidazole, pH of the final buffer should be 7.5). The protein is eluted off of the column with a series of increasing Imidazole solutions made by adjusting the ratios of Lysis Buffer A to Buffer B. Three different concentrations are used: 3 volumes of 75 mM Imidazole, 3 volumes of

150 mM Imidazole, 5 volumes of 500 mM Imidazole. The fractions containing the purified protein are analyzed using 8 %, 10 % or 14% SDS-PAGE depending on the protein size. The purified protein is then dialyzed 2X against phosphate-buffered saline (PBS) in order to place it into an easily workable buffer. The purified protein is stored at 4° C or frozen at -80°.

The following alternative method may be used to purify *B. burgdorferi* expressed in *E coli* when it is present in the form of inclusion bodies. Unless otherwise specified, all of the following steps are conducted at 4-10°C.

Upon completion of the production phase of the *E. coli* fermentation, the cell culture is cooled to 4-10°C and the cells are harvested by continuous centrifugation at 15,000 rpm (Heraeus Sepatech). On the basis of the expected yield of protein per unit weight of cell paste and the amount of purified protein required, an appropriate amount of cell paste, by weight, is suspended in a buffer solution containing 100 mM Tris, 50 mM EDTA, pH 7.4. The cells are dispersed to a homogeneous suspension using a high shear mixer.

The cells are then lysed by passing the solution through a microfluidizer (Microfuidics, Corp. or APV Gaulin, Inc.) twice at 4000-6000 psi. The homogenate is then mixed with NaCl solution to a final concentration of 0.5 M NaCl, followed by centrifugation at 7000 x *g* for 15 min. The resultant pellet is washed again using 0.5M NaCl, 100 mM Tris, 50 mM EDTA, pH 7.4.

The resulting washed inclusion bodies are solubilized with 1.5 M guanidine hydrochloride (GuHCl) for 2-4 hours. After 7000 x *g* centrifugation for 15 min., the pellet is discarded and the *B. burgdorferi* polypeptide-containing supernatant is incubated at 4°C overnight to allow further GuHCl extraction.

Following high speed centrifugation (30,000 x *g*) to remove insoluble particles, the GuHCl solubilized protein is refolded by quickly mixing the GuHCl extract with 20 volumes of buffer containing 50 mM sodium, pH 4.5, 150 mM NaCl, 2 mM EDTA by vigorous stirring. The refolded diluted protein solution is kept at 4°C without mixing for 12 hours prior to further purification steps.

To clarify the refolded *B. burgdorferi* polypeptide solution, a previously prepared tangential filtration unit equipped with 0.16 µm membrane filter with appropriate surface area (e.g., Filtron), equilibrated with 40 mM sodium acetate, pH 6.0 is employed. The filtered sample is loaded onto a cation exchange resin (e.g., Poros HS-50, Perseptive Biosystems). The column is washed with 40 mM sodium acetate, pH 6.0 and eluted with 250 mM, 500 mM, 1000 mM, and 1500 mM NaCl in the same buffer, in a stepwise manner. The absorbance at 280 mm of the effluent is continuously monitored. Fractions are collected and further analyzed by SDS-PAGE.

Fractions containing the *B. burgdorferi* polypeptide are then pooled and mixed with 4

volumes of water. The diluted sample is then loaded onto a previously prepared set of tandem columns of strong anion (Poros HQ-50, Perseptive Biosystems) and weak anion (Poros CM-20, Perseptive Biosystems) exchange resins. The columns are equilibrated with 40 mM sodium acetate, pH 6.0. Both columns are washed with 40 mM sodium acetate, pH 6.0, 200 mM NaCl. The CM-20 column is then eluted using a 10 column volume linear gradient ranging from 0.2 M NaCl, 50 mM sodium acetate, pH 6.0 to 1.0 M NaCl, 50 mM sodium acetate, pH 6.5. Fractions are collected under constant A_{280} monitoring of the effluent. Fractions containing the *B. burgdorferi* polypeptide (determined, for instance, by 16% SDS-PAGE) are then pooled.

The resultant *B. burgdorferi* polypeptide exhibits greater than 95% purity after the above refolding and purification steps. No major contaminant bands are observed from Commassie blue stained 16% SDS-PAGE gel when 5 μg of purified protein is loaded. The purified protein is also tested for endotoxin/LPS contamination, and typically the LPS content is less than 0.1 ng/ml according to LAL assays.

6(b). Alternative Expression and Purification Borrelia polypeptides in E. coli

Tthe vector pQE10 is alternatively used to clone and express some of the polypeptides of the present invention for use in the soft tissue and systemic infection models discussed below. The difference being such that an inserted DNA fragment encoding a polypeptide expresses that polypeptide with the six His residues (i.e., a "6 X His tag") covalently linked to the amino terminus of that polypeptide. The bacterial expression vector pQE10 (QIAGEN, Inc., 9259 Eton Avenue, Chatsworth, CA, 91311) was used in this example . The components of the pQE10 plasmid are arranged such that the inserted DNA sequence encoding a polypeptide of the present invention expresses the polypeptide with the six His residues (*i.e.*, a "6 X His tag")) covalently linked to the amino terminus.

The DNA sequences encoding the desired portions of a polypeptide of SEQ ID NOS:1-155 were amplified using PCR oligonucleotide primers from genomic *B. burgdorferi* DNA. The PCR primers anneal to the nucleotide sequences encoding the desired amino acid sequence of a polypeptide of the present invention. Additional nucleotides containing restriction sites to facilitate cloning in the pQE10 vector were added to the 5' and 3' primer sequences, respectively.

For cloning a polypeptide of the present invention, the 5' and 3' primers were selected to amplify their respective nucleotide coding sequences. One of ordinary skill in the art would appreciate that the point in the protein coding sequence where the 5' and 3' primers begins may be varied to amplify a DNA segment encoding any desired portion of a polypeptide of the present invention. The 5' primer was designed so the coding sequence of the 6 X His tag is aligned with the restriction site so as to maintain its reading frame with that of *B. burgdorferi* polypeptide. The 3' was designed to include an stop codon. The amplified DNA fragment was then cloned, and the protein expressed, as described above for the pQE60 plasmid.

The DNA sequences of SEQ ID NOS:1-155 encoding amino acid sequences may also be cloned and expressed as fusion proteins by a protocol similar to that described directly above, wherein the pET-32b(+) vector (Novagen, 601 Science Drive, Madison, WI 53711) is preferentially used in place of pQE10.

The above methods are not limited to the polypeptide fragements actually produced. The above method, like the methods below, can be used to produce either full length polypeptides or desired fragements therof.

6(c). Alternative Expression and Purification of Borrelia polypeptides in E. coli

The bacterial expression vector pQE60 is used for bacterial expression in this example (QIAGEN, Inc., 9259 Eton Avenue, Chatsworth, CA, 91311). However, in this example, the polypeptide coding sequence is inserted such that translation of the six His codons is prevented and, therefore, the polypeptide is produced with no 6 X His tag.

The DNA sequence encoding the desired portion of the *B. burgdorferi* amino acid sequence is amplified from an *B. burgdorferi* genomic DNA prep the deposited DNA clones using PCR oligonucleotide primers which anneal to the 5' and 3' nucleotide sequences corresponding to the desired portion of the *B. burgdorferi* polypeptides. Additional nucleotides containing restriction sites to facilitate cloning in the pQE60 vector are added to the 5' and 3' primer sequences.

For cloning a *B. burgdorferi* polypeptides of the present invention, 5' and 3' primers are selected to amplify their respective nucleotide coding sequences. One of ordinary skill in the art would appreciate that the point in the protein coding sequence where the 5' and 3' primers begin may be varied to amplify a DNA segment encoding any desired portion of a polypeptide of the present invention. The 3' and 5' primers contain appropriate restriction sites followed by nucleotides complementary to the 5' and 3' ends of the coding sequence respectively. The 3' primer is additionally designed to include an in-frame stop codon.

The amplified *B. burgdorferi* DNA fragments and the vector pQE60 are digested with restriction enzymes recognizing the sites in the primers and the digested DNAs are then ligated together. Insertion of the *B. burgdorferi* DNA into the restricted pQE60 vector places the *B. burgdorferi* protein coding region including its associated stop codon downstream from the IPTG-inducible promoter and in-frame with an initiating AUG. The associated stop codon prevents translation of the six histidine codons downstream of the insertion point.

The ligation mixture is transformed into competent *E. coli* cells using standard procedures such as those described by Sambrook et al. *E. coli* strain M15/rep4, containing multiple copies of the plasmid pREP4, which expresses the lac repressor and confers kanamycin resistance ("Kanr"), is used in carrying out the illustrative example described herein. This strain, which is only one of many that are suitable for expressing *B. burgdorferi* polypeptide, is available commercially (QIAGEN, Inc., *supra*). Transformants are identified by their ability to grow on

LB plates in the presence of ampicillin and kanamycin. Plasmid DNA is isolated from resistant colonies and the identity of the cloned DNA confirmed by restriction analysis, PCR and DNA sequencing.

Clones containing the desired constructs are grown overnight ("O/N") in liquid culture in LB media supplemented with both ampicillin (100 µg/ml) and kanamycin (25 µg/ml). The O/N culture is used to inoculate a large culture, at a dilution of approximately 1:25 to 1:250. The cells are grown to an optical density at 600 nm ("OD600") of between 0.4 and 0.6. isopropyl-b-D-thiogalactopyranoside ("IPTG") is then added to a final concentration of 1 mM to induce transcription from the *lac* repressor sensitive promoter, by inactivating the lacI repressor. Cells subsequently are incubated further for 3 to 4 hours. Cells then are harvested by centrifugation.

To purify the *B. burgdorferi* polypeptide, the cells are then stirred for 3-4 hours at 4°C in 6M guanidine-HCl, pH 8. The cell debris is removed by centrifugation, and the supernatant containing the *B. burgdorferi* polypeptide is dialyzed against 50 mM Na-acetate buffer pH 6, supplemented with 200 mM NaCl. Alternatively, the protein can be successfully refolded by dialyzing it against 500 mM NaCl, 20% glycerol, 25 mM Tris/HCl pH 7.4, containing protease inhibitors. After renaturation the protein can be purified by ion exchange, hydrophobic interaction and size exclusion chromatography. Alternatively, an affinity chromatography step such as an antibody column can be used to obtain pure *B. burgdorferi* polypeptide. The purified protein is stored at 4° C or frozen at -80° C.

The following alternative method may be used to purify *B. burgdorferi* polypeptides expressed in *E coli* when it is present in the form of inclusion bodies. Unless otherwise specified, all of the following steps are conducted at 4-10°C.

Upon completion of the production phase of the *E. coli* fermentation, the cell culture is cooled to 4-10°C and the cells are harvested by continuous centrifugation at 15,000 rpm (Heraeus Sepatech). On the basis of the expected yield of protein per unit weight of cell paste and the amount of purified protein required, an appropriate amount of cell paste, by weight, is suspended in a buffer solution containing 100 mM Tris, 50 mM EDTA, pH 7.4. The cells are dispersed to a homogeneous suspension using a high shear mixer.

The cells ware then lysed by passing the solution through a microfluidizer (Microfuidics, Corp. or APV Gaulin, Inc.) twice at 4000-6000 psi. The homogenate is then mixed with NaCl solution to a final concentration of 0.5 M NaCl, followed by centrifugation at 7000 x *g* for 15 min. The resultant pellet is washed again using 0.5M NaCl, 100 mM Tris, 50 mM EDTA, pH 7.4.

The resulting washed inclusion bodies are solubilized with 1.5 M guanidine hydrochloride (GuHCl) for 2-4 hours. After 7000 x *g* centrifugation for 15 min., the pellet is discarded and the *B. burgdorferi* polypeptide-containing supernatant is incubated at 4°C overnight to allow further GuHCl extraction.

Following high speed centrifugation (30,000 x *g*) to remove insoluble particles, the GuHCl solubilized protein is refolded by quickly mixing the GuHCl extract with 20 volumes of buffer containing 50 mM sodium, pH 4.5, 150 mM NaCl, 2 mM EDTA by vigorous stirring. The refolded diluted protein solution is kept at 4°C without mixing for 12 hours prior to further purification steps.

To clarify the refolded *B. burgdorferi* polypeptide solution, a previously prepared tangential filtration unit equipped with 0.16 µm membrane filter with appropriate surface area (e.g., Filtron), equilibrated with 40 mM sodium acetate, pH 6.0 is employed. The filtered sample is loaded onto a cation exchange resin (e.g., Poros HS-50, Perseptive Biosystems). The column is washed with 40 mM sodium acetate, pH 6.0 and eluted with 250 mM, 500 mM, 1000 mM, and 1500 mM NaCl in the same buffer, in a stepwise manner. The absorbance at 280 mm of the effluent is continuously monitored. Fractions are collected and further analyzed by SDS-PAGE.

Fractions containing the *B. burgdorferi* polypeptide are then pooled and mixed with 4 volumes of water. The diluted sample is then loaded onto a previously prepared set of tandem columns of strong anion (Poros HQ-50, Perseptive Biosystems) and weak anion (Poros CM-20, Perseptive Biosystems) exchange resins. The columns are equilibrated with 40 mM sodium acetate, pH 6.0. Both columns are washed with 40 mM sodium acetate, pH 6.0, 200 mM NaCl. The CM-20 column is then eluted using a 10 column volume linear gradient ranging from 0.2 M NaCl, 50 mM sodium acetate, pH 6.0 to 1.0 M NaCl, 50 mM sodium acetate, pH 6.5. Fractions are collected under constant A_{280} monitoring of the effluent. Fractions containing the *B. burgdorferi* polypeptide (determined, for instance, by 16% SDS-PAGE) are then pooled.

The resultant *B. burgdorferi* polypeptide exhibits greater than 95% purity after the above refolding and purification steps. No major contaminant bands are observed from Commassie blue stained 16% SDS-PAGE gel when 5 µg of purified protein is loaded. The purified protein is also tested for endotoxin/LPS contamination, and typically the LPS content is less than 0.1 ng/ml according to LAL assays.

6(d). Cloning and Expression of B. burgdorferi in Other Bacteria

B. burgdorferi polypeptides can also be produced in: *B. burgdorferi* using the methods of S. Skinner et al., (1988) Mol. Microbiol. 2:289-297 or J. I. Moreno (1996) Protein Expr. Purif. 8(3):332-340; *Lactobacillus* using the methods of C. Rush et al., 1997 Appl. Microbiol. Biotechnol. 47(5):537-542; or in *Bacillus subtilis* using the methods Chang et al., U.S. Patent No. 4,952,508.

7. Cloning and Expression in COS Cells

A *B. burgdorferi* expression plasmid is made by cloning a portion of the DNA encoding a

B. burgdorferi polypeptide into the expression vector pDNAI/Amp or pDNAIII (which can be obtained from Invitrogen, Inc.). The expression vector pDNAI/amp contains: (1) an *E. coli* origin of replication effective for propagation in *E. coli* and other prokaryotic cells; (2) an ampicillin resistance gene for selection of plasmid-containing prokaryotic cells; (3) an SV40 origin of replication for propagation in eukaryotic cells; (4) a CMV promoter, a polylinker, an SV40 intron; (5) several codons encoding a hemagglutinin fragment (i.e., an "HA" tag to facilitate purification) followed by a termination codon and polyadenylation signal arranged so that a DNA can be conveniently placed under expression control of the CMV promoter and operably linked to the SV40 intron and the polyadenylation signal by means of restriction sites in the polylinker. The HA tag corresponds to an epitope derived from the influenza hemagglutinin protein described by Wilson et al. 1984 Cell 37:767. The fusion of the HA tag to the target protein allows easy detection and recovery of the recombinant protein with an antibody that recognizes the HA epitope. pDNAIII contains, in addition, the selectable neomycin marker.

A DNA fragment encoding a *B. burgdorferi* polypeptide is cloned into the polylinker region of the vector so that recombinant protein expression is directed by the CMV promoter. The plasmid construction strategy is as follows. The DNA from a *B. burgdorferi* genomic DNA prep is amplified using primers that contain convenient restriction sites, much as described above for construction of vectors for expression of *B. burgdorferi* in *E. coli*. The 5' primer contains a Kozak sequence, an AUG start codon, and nucleotides of the 5' coding region of the *B. burgdorferi* polypeptide. The 3' primer, contains nucleotides complementary to the 3' coding sequence of the *B. burgdorferi* DNA, a stop codon, and a convenient restriction site.

The PCR amplified DNA fragment and the vector, pDNAI/Amp, are digested with appropriate restriction enzymes and then ligated. The ligation mixture is transformed into an appropriate *E. coli* strain such as SURE™ (Stratagene Cloning Systems, La Jolla, CA 92037), and the transformed culture is plated on ampicillin media plates which then are incubated to allow growth of ampicillin resistant colonies. Plasmid DNA is isolated from resistant colonies and examined by restriction analysis or other means for the presence of the fragment encoding the *B. burgdorferi* polypeptide

For expression of a recombinant *B. burgdorferi* polypeptide, COS cells are transfected with an expression vector, as described above, using DEAE-dextran, as described, for instance, by Sambrook et al. (*supra*). Cells are incubated under conditions for expression of *B. burgdorferi* by the vector.

Expression of the *B. burgdorferi*-HA fusion protein is detected by radiolabeling and immunoprecipitation, using methods described in, for example Harlow et al., *supra*.. To this end, two days after transfection, the cells are labeled by incubation in media containing ^{35}S-cysteine for 8 hours. The cells and the media are collected, and the cells are washed and the lysed with detergent-containing RIPA buffer: 150 mM NaCl, 1% NP-40, 0.1% SDS, 1% NP-40, 0.5% DOC, 50 mM TRIS, pH 7.5, as described by Wilson et al. (*supra*). Proteins are

precipitated from the cell lysate and from the culture media using an HA-specific monoclonal antibody. The precipitated proteins then are analyzed by SDS-PAGE and autoradiography. An expression product of the expected size is seen in the cell lysate, which is not seen in negative controls.

8. Cloning and Expression in CHO Cells

The vector pC4 is used for the expression of *B. burgdorferi* polypeptide in this example. Plasmid pC4 is a derivative of the plasmid pSV2-dhfr (ATCC Accession No. 37146). The plasmid contains the mouse DHFR gene under control of the SV40 early promoter. Chinese hamster ovary cells or other cells lacking dihydrofolate activity that are transfected with these plasmids can be selected by growing the cells in a selective medium (alpha minus MEM, Life Technologies) supplemented with the chemotherapeutic agent methotrexate. The amplification of the DHFR genes in cells resistant to methotrexate (MTX) has been well documented. *See, e.g.*, Alt et al., 1978, J. Biol. Chem. 253:1357-1370; Hamlin et al., 1990, Biochem. et Biophys. Acta, 1097:107-143; Page et al., 1991, Biotechnology 9:64-68. Cells grown in increasing concentrations of MTX develop resistance to the drug by overproducing the target enzyme, DHFR, as a result of amplification of the DHFR gene. If a second gene is linked to the DHFR gene, it is usually co-amplified and over-expressed. It is known in the art that this approach may be used to develop cell lines carrying more than 1,000 copies of the amplified gene(s). Subsequently, when the methotrexate is withdrawn, cell lines are obtained which contain the amplified gene integrated into one or more chromosome(s) of the host cell.

Plasmid pC4 contains the strong promoter of the long terminal repeat (LTR) of the Rouse Sarcoma Virus, for expressing a polypeptide of interest, Cullen, et al. (1985) Mol. Cell. Biol. 5:438-447; plus a fragment isolated from the enhancer of the immediate early gene of human cytomegalovirus (CMV), Boshart, et al., 1985, Cell 41:521-530. Downstream of the promoter are the following single restriction enzyme cleavage sites that allow the integration of the genes: *Bam* HI, *Xba* I, and *Asp* 718. Behind these cloning sites the plasmid contains the 3' intron and polyadenylation site of the rat preproinsulin gene. Other high efficiency promoters can also be used for the expression, e.g., the human ß-actin promoter, the SV40 early or late promoters or the long terminal repeats from other retroviruses, e.g., HIV and HTLVI. Clontech's Tet-Off and Tet-On gene expression systems and similar systems can be used to express the *B. burgdorferi* polypeptide in a regulated way in mammalian cells (Gossen et al., 1992, Proc. Natl. Acad. Sci. USA 89:5547-5551. For the polyadenylation of the mRNA other signals, e.g., from the human growth hormone or globin genes can be used as well. Stable cell lines carrying a gene of interest integrated into the chromosomes can also be selected upon co-transfection with a selectable marker such as gpt, G418 or hygromycin. It is advantageous to use more than one selectable marker in the beginning, e.g., G418 plus methotrexate.

The plasmid pC4 is digested with the restriction enzymes and then dephosphorylated using calf intestinal phosphates by procedures known in the art. The vector is then isolated from

a 1% agarose gel. The DNA sequence encoding the *B. burgdorferi* polypeptide is amplified using PCR oligonucleotide primers corresponding to the 5' and 3' sequences of the desired portion of the gene. A 5' primer containing a restriction site, a Kozak sequence, an AUG start codon, and nucleotides of the 5' coding region of the *B. burgdorferi* polypeptide is synthesized and used. A 3' primer, containing a restriction site, stop codon, and nucleotides complementary to the 3' coding sequence of the *B. burgdorferi* polypeptides is synthesized and used. The amplified fragment is digested with the restriction endonucleases and then purified again on a 1% agarose gel. The isolated fragment and the dephosphorylated vector are then ligated with T4 DNA ligase. *E. coli* HB101 or XL-1 Blue cells are then transformed and bacteria are identified that contain the fragment inserted into plasmid pC4 using, for instance, restriction enzyme analysis.

Chinese hamster ovary cells lacking an active DHFR gene are used for transfection. Five μg of the expression plasmid pC4 is cotransfected with 0.5 μg of the plasmid pSVneo using a lipid-mediated transfection agent such as Lipofectin™ or LipofectAMINE.™ (LifeTechnologies Gaithersburg, MD). The plasmid pSV2-neo contains a dominant selectable marker, the *neo* gene from Tn5 encoding an enzyme that confers resistance to a group of antibiotics including G418. The cells are seeded in alpha minus MEM supplemented with 1 mg/ml G418. After 2 days, the cells are trypsinized and seeded in hybridoma cloning plates (Greiner, Germany) in alpha minus MEM supplemented with 10, 25, or 50 ng/ml of methotrexate plus 1 mg/ml G418. After about 10-14 days single clones are trypsinized and then seeded in 6-well petri dishes or 10 ml flasks using different concentrations of methotrexate (50 nM, 100 nM, 200 nM, 400 nM, 800 nM). Clones growing at the highest concentrations of methotrexate are then transferred to new 6-well plates containing even higher concentrations of methotrexate (1 μM, 2 μM, 5 μM, 10 mM, 20 mM). The same procedure is repeated until clones are obtained which grow at a concentration of 100-200 μM. Expression of the desired gene product is analyzed, for instance, by SDS-PAGE and Western blot or by reversed phase HPLC analysis.

The disclosure of all publications (including patents, patent applications, journal articles, laboratory manuals, books, or other documents) cited herein are hereby incorporated by reference in their entireties.

The present invention is not to be limited in scope by the specific embodiments described herein, which are intended as single illustrations of individual aspects of the invention. Functionally equivalent methods and components are within the scope of the invention, in addition to those shown and described herein and will become apparent to those skilled in the art from the foregoing description and accompanying drawings. Such modifications are intended to fall within the scope of the appended claims.

TABLE 1. Borrelia burgdorferi - Putative coding regions of novel proteins similar to know proteins

Contig ID	ORF ID	Start (nt)	Stop (nt)	match acession	match gene name	% sim	% ident
1	92	100363	100184	gil500722	similar to entire extracellular domain of glycine receptors [Caenorhabditis elegans]	100	66
1	537	513231	513608	gil47453	ribosomal protein S12 [Streptococcus pneumoniae]	92	85
1	283	272186	270849	gil1001376	ATP-dependent protease ATPase subunit [Synechocystis sp.]	89	75
1	847	798835	799131	gil467373	ribosomal protein S18 [Bacillus subtilis]	86	69
1	78	91504	91235	gil1573896	ribosomal protein L27 (rpL27) [Haemophilus influenzae]	85	70
1	732	687538	686753	gil1591672	phosphate transport system ATP-binding protein [Methanococcus jannaschii]	84	65
1	788	739513	739232	gil142459	initiation factor 1 [Bacillus subtilis]	84	68
1	960	901448	901780	gnl|PID|e243769	ORF YGL149w [Saccharomyces cerevisiae]	84	68
1	760	717009	715843	gil623028	orf 361; ranslated orf similarity to SW: RF1_SALTY peptide chain release factor 1 of Salmonella typhimurium [Coxiella burnetii]	83	60
1	115	115536	115312	gil695315	NADH dehydrogenase subunit [Digitalis grandiflora]	82	58
1	184	178954	176918	bbs|57690	EF-G=elongation factor G [Thermotoga maritima, Peptide, 682 aa] [Thermotoga maritima]	82	63
1	447	425980	425453	gil143804	Ndk [Bacillus subtilis]	82	56
1	201	194702	194103	gil530438	arabinose transport protein [Mycoplasma capricolum]	81	53
1	477	446671	445589	gil882454	fructose 1,6-bisphosphate aldolase [Escherichia coli]	81	61
1	601	569453	568650	gil349227	transmembrane protein [Escherichia coli]	81	56
1	887	838084	837224	gil1237019	Srb [Bacillus subtilis]	81	52
1	889	840561	839497	gil154276	peptide chain release factor 2 [Salmonella typhimurium]	81	65
1	896	846681	845440	gil1377823	aminopeptidase [Bacillus subtilis]	81	60
1	60	71604	68890	gil1619909	DNA mismatch repair protein [Thermotoga maritima]	80	59
1	354	348744	349157	gil1765976	chemotaxis protein CheY [Treponema pallidum]	80	42
1	423	409238	408855	gnl|PID|e211829	50S ribosomal protein L14 [Odontella sinensis]	80	61
1	426	410130	409711	gil1652420	50S ribosomal protein L16 [Synechocystis sp.]	80	59
1	507	482736	482936	gil515924	glucosyltransferase [Saccharomyces cerevisiae]	80	40
1	534	505081	505467	pir|A02771|R	ribosomal protein L7/L12 - Micrococcus luteus	80	67

Borrelia burgdorferi - Putative coding regions of novel proteins similar to know proteins

				7MCML			
1	597	567506	566532	gil580899	OppF gene product [Bacillus subtilis]	80	59
1	6	11241	9994	gnl|PID|e242614	arginine deiminase [Clostridium perfringens]	79	62
1	478	447926	446835	gnl|PID|e288124	glucose epimerase [Bacillus thuringiensis]	79	66
1	804	758549	757704	gil455176	glucosamine-6-phosphate deaminase protein [Escherichia coli]	79	60
1	25	31595	31894	gil1017809	similar to dihydropryridine-sensitive l-type, skeletal muscle calcium channel alpha-1 subunit (SP:CIC1_RABIT, P07293) [Caenorhabditis elegans]	78	57
1	134	134667	134323	gil159199	cecropin D [Hyalophora cecropia]	78	50
1	230	215177	216028	gnl|PID|e265537	DnaJ-homologue [Thermus aquaticus thermophilus]	78	59
1	531	503406	503849	gil587583	ribosomal protein L11 [Thermus aquaticus thermophilus]	78	58
1	867	817849	819579	gil912449	Na+ -ATPase alpha subunit [Enterococcus hirae]	78	60
1	127	127383	127745	gil537364	heat shock protein 60 (GroEL) like protein [Porphyromonas gingivalis]	77	60
1	190	182991	182251	gil1235682	mevalonate pyrophosphate decarboxylase [Homo sapiens]	77	51
1	225	213158	212388	gil1651340	Phosphoglycerate mutase 1 [Escherichia coli]	77	59
1	284	272770	272165	gil1001349	ATP-dependent protease ClpP [Synechocystis sp.]	77	62
1	324	318280	314789	gil1573746	DNA polymerase III, alpha chain (dnaE) [Haemophilus influenzae]	77	58
1	555	530150	531370	gil143795	transfer RNA-Tyr synthetase [Bacillus subtilis]	77	52
1	770	722470	722892	gil1653602	hypothetical protein [Synechocystis sp.]	77	54
1	833	790115	790909	gnl|PID|e248886	unknown [Mycobacterium tuberculosis]	77	56
1	52	62205	61918	gnl|PID|e118966	ribosomal protein S15 [Thermus aquaticus thermophilus]	76	60
1	144	141975	141736	bbsl77721	KHS toxin, killer heat sensitive toxin=KHS [Saccharomyces cerevisiae, Peptide, 708 aa] [Saccharomyces cerevisiae]	76	38
1	293	280702	280529	gil1146275	VP2 protein [Bluetongue virus 9]	76	47
1	323	314795	314199	gil1651915	hypothetical protein [Synechocystis sp.]	76	48
1	362	356749	355508	gil633147	ribose-phosphate pyrophosphokinase [Bacillus caldolyticus]	76	44

Borrelia burgdorferi - Putative coding regions of novel proteins similar to know proteins

1	410	403332	402922	gil606232	30S ribosomal subunit protein S11 [Escherichia coli]	76	52
1	411	403754	403341	gil1652405	30S ribosomal protein S13 [Synechocystis sp.]	76	55
1	454	431743	431003	gil1016012	neural cell adhesion protein BIG-2 precursor [Rattus norvegicus]	76	61
1	710	670457	671569	gil467376	unknown [Bacillus subtilis]	76	58
1	873	824849	826675	gil1303804	YqeQ [Bacillus subtilis]	76	52
1	942	886017	886751	gil1183839	unknown [Pseudomonas aeruginosa]	76	54
1	5	9956	8943	gil1552842	OTCase [Escherichia coli]	75	62
1	51	61909	59735	gil1184680	polynucleotide phosphorylase [Bacillus subtilis]	75	54
1	55	66283	63620	gil39954	IF2 (aa 1-741) [Bacillus stearothermophilus]	75	53
1	83	93454	94410	gnl|PID|e289138	similar to flagellar hook-basal body proteins [Bacillus subtilis]	75	46
1	88	97435	98283	gil687583	RpoS [Yersinia enterocolitica]	75	47
1	245	229112	230158	gil1574806	spermidine/putrescine transport ATP-binding protein (potA) [Haemophilus influenzae]	75	55
1	264	251076	250801	gil1763634	alpha1A-voltage-dependent calcium channel [Homo sapiens]	75	60
1	297	285723	284461	gil556886	serine hydroxymethyltransferase [Bacillus subtilis]	75	58
1	375	367682	366903	gil467372	3'-exo-deoxyribonuclease [Bacillus subtilis]	75	62
1	385	378055	377114	gil45986	NAD synthetase [Rhodobacter capsulatus]	75	55
1	416	406437	405925	gil1044981	ribosomal protein S5 [Bacillus subtilis]	75	56
1	418	407390	406812	gil600032	L6 ribosomal protein [Streptomyces coelicolor]	75	53
1	424	409520	409251	gil44218	ribosomal protein S17 (AA 1-85) [Mycoplasma capricolum]	75	58
1	530	502806	503366	gil396321	nusG [Escherichia coli]	75	56
1	548	523428	522904	gil1573470	H. influenzae predicted coding region HI0491 [Haemophilus influenzae]	75	55
1	575	546579	548393	pirlC30010lC30010	hypothetical ORF-6 protein - Sauroleishmania tarentolae mitochondrion (SGC6)	75	50
1	906	854433	855215	gil511148	hemolysin [Serpulina hyodysenteriae]	75	56
1	68	85054	83102	gil467458	cell division protein [Bacillus subtilis]	74	57
1	162	158608	157502	gil531460	Mbl protein [Bacillus subtilis]	74	49
1	177	172327	171950	pirlA45434lA45434	ribosomal protein L19 - Bacillus stearothermophilus	74	54
1	475	443773	445203	gil396501	aspartyl-tRNA synthetase [Thermus aquaticus thermophilus]	74	52

Borrelia burgdorferi - Putative coding regions of novel proteins similar to know proteins

1	549	524561	523374	gil1020317	S-adenosylmethionine synthetase [Staphylococcus aureus]	74	57
1	595	565672	564347	gil460259	enolase [Bacillus subtilis]	74	58
1	720	681529	680489	gil1651962	hypothetical protein [Synechocystis sp.]	74	49
1	745	702297	701173	gil289287	UDP-glucose pyrophosphorylase [Bacillus subtilis]	74	50
1	13	20409	17551	gil1652531	excinuclease ABC subunit A [Synechocystis sp.]	73	56
1	98	103790	104947	gil514330	sensor kinase [Bacillus subtilis]	73	49
1	188	182064	181102	gil887601	Erg8p [Saccharomyces cerevisiae]	73	43
1	314	303616	302786	gil473817	'ORF' [Escherichia coli]	73	53
1	366	358916	361078	gnl|PID|e245791	ORF YLR069c [Saccharomyces cerevisiae]	73	51
1	444	424047	423181	gil1574704	hypothetical [Haemophilus influenzae]	73	51
1	556	531372	533672	gil511145	hemolysin [Serpulina hyodysenteriae]	73	52
1	576	548257	548045	gil406135	glycoprotein 120 [Simian immunodeficiency virus]	73	53
1	598	568379	567504	gil143607	sporulation protein [Bacillus subtilis]	73	55
1	604	572375	570729	bbsl161785	60 kda antigen [Borrelia coriaceae, C053, ATCC 4338, Peptide, 514 aa] [Borrelia coriaceae]	73	53
1	674	634175	633648	gil1595810	type-I signal peptidase SpsB [Staphylococcus aureus]	73	47
1	692	654267	651727	gnl|PID|e268456	unknown [Mycobacterium tuberculosis]	73	54
1	719	679186	680499	gil500705	Similar to Seryl-tRNA synthetase [Saccharomyces cerevisiae]	73	56
1	725	682189	682899	gnl|PID|e243681	ORF YGR248w [Saccharomyces cerevisiae]	73	63
1	895	845455	844964	gil1652288	hypothetical protein [Synechocystis sp.]	73	50
1	16	24242	26497	gil511145	hemolysin [Serpulina hyodysenteriae]	72	53
1	99	104935	106305	gil619917	NtrC/NifA-like protein regulator [Escherichia coli]	72	54
1	133	134036	135055	gil556881	Similar to Saccharomyces cerevisiae SUA5 protein [Bacillus subtilis]	72	51
1	270	256925	260308	gil467444	transcription-repair coupling factor [Bacillus subtilis]	72	49
1	280	267529	268221	gil1573812	ribosomal protein S4 (rpS4) [Haemophilus influenzae]	72	51
1	282	270922	268472	gil402504	lon protease [Bacillus brevis]	72	51
1	325	319544	318363	gil48362	haemolysin releasing protein (AA 1-548) [Vibrio cholerae]	72	41
1	328	322678	321053	gil1591801	CTP synthase [Methanococcus jannaschii]	72	42

Borrelia burgdorferi - Putative coding regions of novel proteins similar to know proteins

1	348	341460	341182	gil145687	ptsH protein [Escherichia coli]	72	55
1	405	399941	399096	gil1045937	M. genitalium predicted coding region MG246 [Mycoplasma genitalium]	72	53
1	420	408009	407779	gil580930	S14 protein (AA 1-61) [Bacillus subtilis]	72	55
1	593	563383	563850	gil1574283	ribosomal protein L13 (rpL13) [Haemophilus influenzae]	72	54
1	682	641030	643399	gil1574437	sporulation protein (spoIIIE) [Haemophilus influenzae]	72	51
1	754	710160	710750	gil460080	D-alanine:D-alanine ligase-related protein [Enterococcus faecalis]	72	47
1	767	721422	721640	gil868029	large ribosomal subunit protein L35 [Buchnera aphidicola]	72	48
1	860	811923	810511	gil1001357	asparaginyl-tRNA synthetase [Synechocystis sp.]	72	54
1	14	22434	20407	gil1737482	UvrB [Helicobacter pylori]	71	52
1	72	87471	87674	gil1016781	beta-b protein [Barley stripe mosaic virus]	71	42
1	289	278760	278239	gil534842	ORF9 [Rhizobium meliloti]	71	43
1	307	298685	296736	gil1652099	long-chain-fatty-acid CoA ligase [Synechocystis sp.]	71	48
1	321	313551	312130	gil1732243	RecG [Treponema pallidum]	71	52
1	522	494911	496383	gil459009	similar to multifunctional aminoacyl-tRNA synthetase, especially to the prolyl-tRNA synthetase region [Caenorhabditis elegans]	71	48
1	554	528795	530156	pirlS58522lS58522	glycyl-tRNA synthetase - Thermus thermophilus	71	54
1	582	553725	552271	gil285623	pyruvate kinase [Bacillus stearothermophilus]	71	52
1	684	644626	643661	gil217121	ORF1 [Synechococcus elongatus]	71	52
1	723	681731	681561	gil44228	secretion protein SecY (AA 1-482) [Mycoplasma capricolum]	71	42
1	856	806939	807700	gil216341	ORF for methionine amino peptidase [Bacillus subtilis]	71	53
1	947	890096	890665	gil147485	queA [Escherichia coli]	71	56
1	28	38112	40613	gil1439562	Cdc28p [Schizosaccharomyces pombe]	70	53
1	36	45750	44806	gil290494	o287 [Escherichia coli]	70	32
1	84	94408	95220	gil47677	flgG protein product (AA 1-260) [Salmonella typhimurium]	70	50
1	128	127889	128569	gil1574387	H. influenzae predicted coding region HI1534 [Haemophilus influenzae]	70	58
1	468	441049	441330	gil1673757	(AE000012) Mycoplasma pneumoniae, phosphocarrier protein HPr; similar to GenBank Accession Number A49683, from M. capricolum [Mycoplasma pneumoniae]	70	41
1	532	503834	504529	splQ06797lRL1_BACSU	50S RIBOSOMAL PROTEIN L1 (BL1).	70	48

Borrelia burgdorferi - Putative coding regions of novel proteins similar to know proteins

1	594	563858	564280	gil606169	30S ribosomal subunit protein S9 [Escherichia coli]	70	56
1	622	591070	591606	gil153906	CheW protein [Salmonella typhimurium]	70	48
1	703	664161	662611	gnl|PID|e283919	glycerol kinase [Sulfolobus solfataricus]	70	60
1	726	682886	682659	gil836815	cdc4 gene product which is essential for initiation of DNA replication in yeast [Saccharomyces cerevisiae]	70	35
1	766	720854	721417	gil436165	Dsg [Myxococcus xanthus]	70	47
1	768	721649	722008	gnl|PID|e254981	ribosomal protein L20 [Bacillus subtilis]	70	48
1	965	904395	905465	gil1100074	tryptophanyl-tRNA synthetase [Clostridium longisporum]	70	47
1	87	96986	97336	gil160092	asparagine-rich protein [Plasmodium falciparum]	69	46
1	110	112658	113602	gil1001733	ABC transporter [Synechocystis sp.]	69	46
1	181	174037	173762	pir||C47154|C47154	ribosomal protein S16 - Bacillus subtilis	69	52
1	233	219872	218076	gil1001493	protein-export membrane protein SecD [Synechocystis sp.]	69	47
1	234	220245	219922	gil1402532	ORF11 [Enterococcus faecalis]	69	32
1	373	366148	363977	gil1574200	hypothetical [Haemophilus influenzae]	69	48
1	419	407781	407371	gil498771	ribosomal S8 protein [Thermus aquaticus thermophilus]	69	46
1	517	489315	491207	gil151932	fructose enzyme II [Rhodobacter capsulatus]	69	42
1	600	568891	568388	gil143606	sporulation protein [Bacillus subtilis]	69	44
1	733	689098	687536	gil1303856	YqgI [Bacillus subtilis]	69	46
1	874	826778	827746	pir||S08183|S08183	L-lactate dehydrogenase (EC 1.1.1.27) X - Bacillus psychrosaccharolyticus	69	50
1	894	844392	844547	gil1592324	M. jannaschii predicted coding region MJ1172 [Methanococcus jannaschii]	69	53
1	934	879725	879237	gil153566	ORF (19K protein) [Enterococcus faecalis]	69	42
1	49	57779	57976	gil809583	unknown [Saccharomyces cerevisiae]	68	36
1	107	110374	111513	gnl|PID|e255943	M04B2.4 [Caenorhabditis elegans]	68	48
1	132	133978	133148	gil1001663	rare lipoprotein A [Synechocystis sp.]	68	53
1	142	141239	142642	gnl|PID|e233874	hypothetical protein [Bacillus subtilis]	68	45
1	148	145381	144005	gil558574	pyrophosphate--fructose-6-phosphate 1-phosphotransferase	68	48

Borrelia burgdorferi - Putative coding regions of novel proteins similar to know proteins

					[Entamoeba histolytica]		
1	151	147107	147295	gil193442	beta-galactoside binding protein [Mus musculus]	68	42
1	173	170051	169296	gil1389549	ORF3 [Bacillus subtilis]	68	50
1	182	175384	174035	gil1653392	signal recognition particle protein [Synechocystis sp.]	68	45
1	203	195720	195439	gil146854	galactose binding protein [Escherichia coli]	68	42
1	243	228084	227344	gil1573101	hypothetical [Haemophilus influenzae]	68	41
1	255	237406	235943	gil1652530	glucose 6-phosphate dehydrogenase [Synechocystis sp.]	68	47
1	263	249185	251077	gil1573422	fructose-permease IIBC component (fruA) [Haemophilus influenzae]	68	42
1	372	364016	363333	gil976104,	MHc, class IIB gene product [Poecilia reticulata]	68	50
1	407	401421	399874	gil150209	ORF 1 [Mycoplasma mycoides]	68	40
1	412	405181	403862	gil699588	SecY protein [Corynebacterium glutamicum]	68	44
1	425	409759	409508	gil1591164	ribosomal protein L29 [Methanococcus jannaschii]	68	51
1	506	482737	482087	gil1498049	adenylate kinase [Paracoccus denitrificans]	68	37
1	550	525298	524549	gil473817	'ORF' [Escherichia coli]	68	42
1	697	658281	657673	gnl|PID|e255117	hypothetical protein [Bacillus subtilis]	68	51
1	717	678378	675460	gil927711	D9461.18p; CAI: 0.15 [Saccharomyces cerevisiae]	68	52
1	730	685905	686750	gil1707057	coded for by C. elegans cDNA CEESS55F; coded for by C. elegans cDNA yk84a1.3; coded for by C. elegans cDNA yk78g7.3; coded for by C. elegans cDNA yk168g9.5; coded for by C. elegans cDNA yk78g7.5; coded for by C. elegans cDNA yk84a1.5; strong s	68	52
1	747	704201	702609	gil466482	outer surface protein F [Borrelia burgdorferi]	68	44
1	793	744799	745815	gil1652037	hypothetical protein [Synechocystis sp.]	68	50
1	843	796077	797807	gil1072418	glcA gene product [Staphylococcus carnosus]	68	50
1	875	829335	827776	gil147774	ORF III [Escherichia coli]	68	47
1	900	850141	850680	gnl|PID|e290282	polypeptide deformylase [Calothrix PCC7601]	68	52
1	15	22531	24255	gnl|PID|e280490	unknown [Streptococcus pneumoniae]	67	43
1	20	26452	29640	gil498991	ILS1 protein [Saccharomyces cerevisiae]	67	49
1	27	35545	38139	gil1652946	DNA topoisomerase I [Synechocystis sp.]	67	47

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Borrelia burgdorferi - Putative coding regions of novel proteins similar to know proteins

1	111	113572	114333	gil1001529	hypothetical protein [Synechocystis sp.]	67	36
1	170	166286	165876	gil567036	CapE [Staphylococcus aureus]	67	35
1	202	195499	194651	gil1674275	(AE000056) Mycoplasma pneumoniae, hypothetical ABC transporter (yjcW) homolog; similar to Swiss-Prot Accession Number P32721, from E. coli [Mycoplasma pneumoniae]	67	41
1	206	197487	197098	gil1653841	P protein [Synechocystis sp.]	67	35
1	271	260292	261551	gil349834	acetate kinase [Methanosarcina thermophila]	67	44
1	313	302731	301643	gnl|PID|e249981	phosphotransacetylase [Thermoanaerobacterium thermosaccharolyticum]	67	51
1	422	408897	408535	pir|A02819|R5BS24	ribosomal protein L24 - Bacillus stearothermophilus	67	49
1	480	450326	448689	gil1574032	hypothetical [Haemophilus influenzae]	67	42
1	529	502315	502509	gil1001264	50S ribosomal protein L33 [Synechocystis sp.]	67	56
1	588	559618	561111	gil1224069	amidase [Moraxella catarrhalis]	67	51
1	683	643676	643437	gil710340	ribosomal protein S21 [Myxococcus xanthus]	67	49
1	698	658454	659500	gil460955	TagE [Vibrio cholerae]	67	38
1	700	660039	660536	gil467420	unknown [Bacillus subtilis]	67	42
1	729	684089	685888	gnl|PID|e267607	alanyl-tRNA synthatase [Thermus aquaticus thermophilus]	67	51
1	835	791754	792341	gnl|PID|e248763	unknown [Mycobacterium tuberculosis]	67	46
1	857	807722	809191	gil1526428	GsrA protein [Yersinia enterocolitica]	67	46
1	868	819577	820905	gil1590954	ATP synthase, subunit B [Methanococcus jannaschii]	67	53
1	74	88393	88028	gil1572979	hypothetical [Haemophilus influenzae]	66	43
1	91	99152	100252	gil561690	sialoglycoprotease [Pasteurella haemolytica]	66	44
1	123	121472	120783	gil1652843	endonuclease III [Synechocystis sp.]	66	42
1	149	146362	145379	gil1216385	orf304 gene product [Treponema pallidum]	66	43
1	185	179585	179001	gil1574811	neutrophil activating protein (napA) [Haemophilus influenzae]	66	49
1	275	265075	265584	gil401785	cytidine deaminase [Mycoplasma pirum]	66	41
1	330	324514	323696	gil1574641	ribonucleotide transport ATP-binding protein (mkl) [Haemophilus influenzae]	66	41
1	335	327265	326888	gil510670	cheY gene product [Rhodobacter sphaeroides]	66	44
1	355	349142	349603	gil499382	FliS [Bacillus subtilis]	66	28

Borrelia burgdorferi - Putative coding regions of novel proteins similar to know proteins

1	358	351051	350827	gil1546788	tar-1 [Trichostrongylus colubriformis]	66	55
1	404	399121	398324	gil296626	hemolysin [Serpulina hyodysenteriae]	66	53
1	491	461335	460550	gil45713	P.putida genes rpmH, rnpA, 9k, 60k, 50k, gidA, gidB, uncI and uncB [Pseudomonas putida]	66	41
1	513	486046	485159	gil153903	methyltransferase (cheR; EC 2.1.1.24) [Salmonella typhimurium]	66	42
1	552	526495	527316	gil340613	A 'c' was inserted after nt 369 (=nt 10459 in genomic sequence (M10126)) to correct -1 frameshift probably due to gel compression [Leishmania tarentolae]	66	40
1	611	579933	581069	gil886130	putative pectinesterase [Medicago sativa]	66	33
1	627	595395	596288	gnl|PID|e2639 31	OrfD [Streptococcus pneumoniae]	66	47
1	772	723788	723522	gil1762342	could accelerate degradation of certain transcripts [Bacillus subtilis]	66	47
1	816	770251	770060	gil393266	glycerol ester hydrolase [Staphylococcus aureus]	66	33
1	841	795927	795208	gil662880	novel hemolytic factor [Bacillus cereus]	66	46
1	882	835002	834262	gil862629	similar to the ATP-binding transport protein family [Buchnera aphidicola]	66	40
1	73	87915	87619	gil39656	spoVG gene product [Bacillus megaterium]	65	40
1	97	103039	102803	gil532272	phosphatidylserine decarboxylase [Bacillus subtilis]	65	39
1	106	110281	109649	gil1377852	ClpP [Yersinia enterocolitica]	65	42
1	159	156186	154372	gil1572977	penicillin-binding protein 2 (pbp2) [Haemophilus influenzae]	65	41
1	172	168084	169325	gil1146238	poly(A) polymerase [Bacillus subtilis]	65	38
1	268	255918	253819	gil829194	bacterial cell wall hydrolase [Enterococcus faecalis]	65	43
1	353	348568	346553	gil1574651	DNA ligase (lig) [Haemophilus influenzae]	65	45
1	696	657577	655781	gil1651216	Pz-peptidase [Bacillus licheniformis]	65	47
1	741	695297	693456	gil1575784	DNA mismatch repair protein [Aquifex pyrophilus]	65	45
1	846	798339	798827	gil1001362	single-stranded DNA-binding protein [Synechocystis sp.]	65	45
1	932	876643	878559	gil508471	gyrase A [Helicobacter pylori]	65	40
1	936	881238	882224	gil1652260	leader peptidase I [Synechocystis sp.]	65	40
1	961	902331	901519	gil1256146	YbbQ [Bacillus subtilis]	65	48
1	963	903280	904407	gil1573307	hypothetical [Haemophilus influenzae]	65	41
1	37	47101	45683	gil556014	UDP-N-acetyl muramate-alanine ligase [Bacillus subtilis]	64	46

Borrelia burgdorferi - Putative coding regions of novel proteins similar to know proteins

1	61	72211	71642	gil1041785	rhoptry protein [Plasmodium yoelii]	64	41
1	130	131969	129336	gil1574225	valyl-tRNA synthetase (valS) [Haemophilus influenzae]	64	43
1	156	152924	151140	gil43066	threonyl-tRNA synthetase (thrS; EC 6.1.1.3) [Escherichia coli]	64	43
1	174	170326	170033	gil1652390	acyl carrier protein [Synechocystis sp.]	64	48
1	175	171105	170545	gil1573650	lipopolysaccharide core biosynthesis protein (kdtB) [Haemophilus influenzae]	64	45
1	178	173033	172293	gil1046163	tRNA (guanine-N1)-methyltransferase [Mycoplasma genitalium]	64	47
1	180	173764	173513	gnl|PID|e248893	unknown [Mycobacterium tuberculosis]	64	34
1	207	197654	197436	gil11665	ORF2136 [Marchantia polymorpha]	64	47
1	217	206795	205761	gil1652866	N-acetylmuramoyl-L-alanine amidase [Synechocystis sp.]	64	30
1	244	228146	229036	gil1046160	hypothetical protein (GB:U00021_5) [Mycoplasma genitalium]	64	33
1	246	230149	230967	gil147336	transmembrane protein [Escherichia coli]	64	36
1	267	253160	253723	gil467430	unknown [Bacillus subtilis]	64	52
1	340	333349	332783	gil145520	cheW peptide [Escherichia coli]	64	42
1	384	376509	375565	gil1653737	monophosphatase [Synechocystis sp.]	64	44
1	449	428137	426437	gil467409	DNA polymerase III subunit [Bacillus subtilis]	64	41
1	510	484558	483998	pir||A00547|XYEBET	protein-glutamate methylesterase (EC 3.1.1.61) - Salmonella typhimurium	64	45
1	603	570416	569451	gil1574678	dipeptide transport system permease protein (dppB) [Haemophilus influenzae]	64	44
1	679	637996	640224	gil1001335	soluble lytic transglycosylase [Synechocystis sp.]	64	42
1	753	709637	710194	gnl|PID|e283360	unknown [Mycobacterium tuberculosis]	64	42
1	817	771784	771969	gnl|PID|e250307	W04B2.3 [Caenorhabditis elegans]	64	41
1	839	793892	795211	gil1573939	hypothetical [Haemophilus influenzae]	64	38
1	861	811972	812853	gil396314	glutamate synthase [Escherichia coli]	64	38
1	870	821501	823339	gil472918	v-type Na-ATPase [Enterococcus hirae]	64	42
1	901	850668	851615	gil581088	methionyl-tRNA formyltransferase [Escherichia coli]	64	38
1	904	853492	853884	gil992960	thioredoxin [Arabidopsis thaliana]	64	41
1	24	34314	31444	gil42914	SbcC (AA 1-1048) [Escherichia coli]	63	45

Borrelia burgdorferi - Putative coding regions of novel proteins similar to know proteins

1	77	91198	90194	gil1652022	GTP-binding protein [Synechocystis sp.]	63	45
1	209	198041	197862	gil11665	ORF2136 [Marchantia polymorpha]	63	42
1	227	214639	213956	gil1652349	oxygen independent coprophorphyrinogen III oxidase [Synechocystis sp.]	63	41
1	232	218116	217193	gil1573204	protein-export membrane protein (secF) [Haemophilus influenzae]	63	36
1	247	230965	231762	spIP45169IPOTC_HAEIN	SPERMIDINE/PUTRESCINE TRANSPORT SYSTEM PERMEASE PROTEIN POTC.	63	36
1	272	262171	261614	gil1780755	DJ-1 protein [Homo sapiens]	63	46
1	290	279964	278735	gil143439	DD-carboxypeptidase [Bacillus subtilis]	63	41
1	333	326012	325818	gil293954,	mating type a-1 protein [Neurospora crassa]	63	42
1	508	484000	482759	gil1041116	TRAB [Plasmid pPD1]	63	34
1	553	527314	528801	spIP15189ISYE_RHIME	GLUTAMYL-TRNA SYNTHETASE (EC 6.1.1.17) (GLUTAMATE--TRNA LIGASE) (GLURS).	63	48
1	569	542317	543747	gil1652577	carboxyl-terminal protease [Synechocystis sp.]	63	49
1	620	590442	589360	gil1098641	Bts1p [Saccharomyces cerevisiae]	63	43
1	701	660784	660623	gil1339938	EC 1.1.99.5 [Mus musculus]	63	47
1	702	662231	660735	gil763191	glycerol 3 phosphate dehydrogenase [Saccharomyces cerevisiae]	63	37
1	704	664938	664159	gil142997	glycerol uptake facilitator [Bacillus subtilis]	63	45
1	746	702035	702631	gil41497	ORF 4 (AA 1-198); 20 kD [Escherichia coli]	63	36
1	748	705645	704671	gil436158	putative integral membrane protease required for high frequency lysogenization by bacteriophage lambda [Escherichia coli]	63	33
1	749	706431	705643	gil507734	HflK [Vibrio parahaemolyticus]	63	28
1	756	715040	713019	gil407881	stringent response-like protein [Streptococcus equisimilis]	63	40
1	825	780572	783289	gil746399	transcription elongation factor [Escherichia coli]	63	43
1	853	803786	804832	gil155055	basic membrane protein precursor [Treponema pallidum]	63	36
1	4	8945	7467	gil1573583	H. influenzae predicted coding region HI0594 [Haemophilus influenzae]	62	38
1	42	50587	51786	gil1573978	pantothenate metabolism flavoprotein (dfp) [Haemophilus influenzae]	62	41
1	57	67740	66271	gil49316	ORF2 gene product [Bacillus subtilis]	62	36
1	64	78979	79767	gil1001473	hypothetical protein [Synechocystis sp.]	62	48
1	80	92123	91806	gnlIPIDle211848	50S ribosomal protein L21 [Odontella sinensis]	62	34

Borrelia burgdorferi - Putative coding regions of novel proteins similar to know proteins

1	101	107458	106793	gil1652679	hypothetical protein [Synechocystis sp.]	62	34
1	102	107464	107883	gil15893	predicted 12.5Kd protein [Mycobacteriophage I5]	62	44
1	226	213238	213969	gil1001678	ribose 5-phosphate isomerase [Synechocystis sp.]	62	39
1	266	251889	253175	gil529118	similar to APE1/LAP4, vacuolar aminopeptidase [Saccharomyces cerevisiae]	62	42
1	299	288749	287274	gil289284	cysteinyl-tRNA synthetase [Bacillus subtilis]	62	46
1	357	349982	352714	gil1633576	similar to proofreading 3'-5' exonuclease and polymerase [Treponema pallidum]	62	41
1	443	423190	422495	gil312380	putative orfW gene product [Clostridium acetobutylicum]	62	32
1	489	458740	459582	gil40031	spoOJ93 gene product [Bacillus subtilis]	62	36
1	511	485147	484494	gil145524	cheB peptide [Escherichia coli]	62	28
1	518	491201	492322	gil146722	phosphomannose isomerase [Escherichia coli]	62	45
1	685	646727	644598	gil1574144	single-stranded-DNA-specific exonuclease (recJ) [Haemophilus influenzae]	62	40
1	695	655800	655063	gil1477770	unknown [Helicobacter pylori]	62	37
1	758	715668	714979	gil1574130	protoporphyrinogen oxidase (hemK) [Haemophilus influenzae]	62	36
1	762	718374	719198	gil1652444	hypothetical protein [Synechocystis sp.]	62	40
1	837	792941	793891	gil1652668	phosphatidate cytidylyltransferase [Synechocystis sp.]	62	41
1	917	862498	862737	gil440851	collagenase [Clostridium perfringens]	62	29
1	46	55889	54726	gil498141	tRNA guanine transglycosylase [Zymomonas mobilis]	61	35
1	81	92710	92174	gil726305	adenine phosphoribosyltransferase form 1 [Triticum aestivum]	61	45
1	100	106820	106557	gil460955	TagE [Vibrio cholerae]	61	45
1	109	111699	112664	gil1001126	hypothetical protein [Synechocystis sp.]	61	48
1	157	154445	153051	gil143657	endospore forming protein [Bacillus subtilis]	61	40
1	193	185315	184227	gil148409	gene not found in Erwinia uredovora crt gene cluster; ORF6 [Erwinia herbicola]	61	42
1	223	209790	210668	splP37214lERA_STRMU	GTP-BINDING PROTEIN ERA HOMOLOG.	61	37
1	273	262392	264062	gil438455	possible N-terminal signal sequence; mature protein may be membrane-anchored and start at Cys-17. 17.5% identity over 354-aa overlap with Candida pelliculosa beta-glucosidase.; putative [Bacillus subtilis]	61	37
1	277	265982	265581	gil1513240	ORFveg110 [Dictyostelium discoideum]	61	29

Borrelia burgdorferi - Putative coding regions of novel proteins similar to know proteins

1	301	291935	289686	gil1354776	MCP-1 [Treponema pallidum]	61	43
1	322	314201	313338	gil1732243	RecG [Treponema pallidum]	61	37
1	380	371430	372392	gil973332	OrfC [Bacillus subtilis]	61	38
1	408	401874	401479	gil147716	ribosomal protein L17 [Escherichia coli]	61	44
1	413	404277	404444	gil1185286	ORF [Sulfolobus shibatae]	61	47
1	415	405927	405616	pirlA02827lR5BS3F	ribosomal protein L30 - Bacillus stearothermophilus	61	31
1	417	406848	406435	pirlB29102lR5BS8F	ribosomal protein L18 - Bacillus stearothermophilus	61	44
1	441	421784	421224	gil153045.	prolipoprotein signal peptidase [Staphylococcus aureus]	61	29
1	467	440722	441042	gil173128	ubiquitin-specific processing protease [Saccharomyces cerevisiae]	61	32
1	613	582695	581547	gil1303756	YqbP [Bacillus subtilis]	61	38
1	615	584397	585476	gil551522	TpN38(b) [Treponema pallidum]	61	26
1	673	632123	633622	gil143999	dnaK homologue [Borrelia burgdorferi]	61	41
1	675	634207	635469	gil1653709	lipoprotein NlpD [Synechocystis sp.]	61	50
1	743	699438	698647	gil1303863	YqgP [Bacillus subtilis]	61	45
1	897	847575	846688	gil1573586	hydrolase (GB:Z33006_1) [Haemophilus influenzae]	61	43
1	938	882836	883282	gil1303831	YqfM [Bacillus subtilis]	61	36
1	7	10415	10627	gnllPIDle211990	T24A11.1 [Caenorhabditis elegans]	60	45
1	23	31428	30475	gil1303865	YqgR [Bacillus subtilis]	60	45
1	35	44812	44267	gil1591369	cytidylate kinase [Methanococcus jannaschii]	60	49
1	198	192994	192053	gil1045801	hypothetical protein (SP:P32720) [Mycoplasma genitalium]	60	33
1	347	341167	339440	gil602680	phosphocarrier protein (enzyme I) [Mycoplasma capricolum]	60	37
1	369	361817	362233	gil1372995	OrfH [Borrelia burgdorferi]	60	37
1	409	402924	401872	gil142463	RNA polymerase alpha-core-subunit [Bacillus subtilis]	60	40
1	438	420142	418793	gnllPIDle276830	UDP-N-acetylglucosamine 1-carboxyvinyltransferase [Bacillus subtilis]	60	40
1	566	540696	539698	gil1573923	prolipoprotein diacylglyceryl transferase (lgt) [Haemophilus influenzae]	60	57
1	587	559368	559655	gil1335805	CD45 homolog [Heterodontus francisci]	60	26
1	589	561098	562558	gil1653395	PET112 [Synechocystis sp.]	60	37

Borrelia burgdorferi - Putative coding regions of novel proteins similar to know proteins

1	736	690957	690076	gil1001260	hypothetical protein [Synechocystis sp.]	60	47
1	738	691078	691659	gil1399829	elongation factor P [Synechococcus PCC7942]	60	34
1	750	707879	706626	gil1573060	hypothetical [Haemophilus influenzae]	60	33
1	784	734589	735635	gil1164996	mxaC gene product [Methylobacterium extorquens]	60	26
1	829	785899	786567	gil1046033	cytidylate kinase [Mycoplasma genitalium]	60	38
1	862	812835	813773	gil1574569	hypothetical [Haemophilus influenzae]	60	36
1	863	813727	816105	gnl|PID|e255093	hypothetical protein [Bacillus subtilis]	60	38
1	878	831250	829943	gil1742766	NifS protein. [Escherichia coli]	60	34
1	929	872578	874110	gil1002666	unknown [Schistosoma mansoni]	60	30
1	937	882211	882861	gil1595810	type-I signal peptidase SpsB [Staphylococcus aureus]	60	40
1	54	63629	63234	gil580902	ORF6 gene product [Bacillus subtilis]	59	38
1	96	102744	103802	gil467409	DNA polymerase III subunit [Bacillus subtilis]	59	40
1	120	118925	119914	gil1574678	dipeptide transport system permease protein (dppB) [Haemophilus influenzae]	59	33
1	140	139567	141174	gil42377	phosphoglucose isomerase (AA 1-549) [Escherichia coli]	59	42
1	195	186577	187659	gil1573129	hypothetical [Haemophilus influenzae]	59	41
1	259	242174	245713	gil1574781	exodeoxyribonuclease V (recB) [Haemophilus influenzae]	59	38
1	288	278281	276257	pir||D64084|D64084	rep helicase, single-stranded DNA-dependent ATPase (rep) homolog - Haemophilus influenzae (strain Rd KW20)	59	36
1	291	280005	281525	gil882504	ORF_f560 [Escherichia coli]	59	34
1	306	294923	296707	gil487937	Similar to arginyl-tRNA synthetase (E. coli) [Saccharomyces cerevisiae]	59	35
1	332	325664	324564	gil466753	alternate gene name yibD [Escherichia coli]	59	39
1	414	405646	405179	gil216338	ORF for L15 ribosomal protein [Bacillus subtilis]	59	40
1	465	439470	440759	gil39269	sigma factor (ntrA) (AA 1-502) [Azotobacter vinelandii]	59	35
1	492	462064	461411	pir||A30191|A30191	hypothetical protein L - Bacillus subtilis (fragment)	59	39
1	495	462955	463752	gil467425	unknown [Bacillus subtilis]	59	38
1	503	480078	481016	gil1651878	regulatory components of sensory transduction system [Synechocystis sp.]	59	38
1	523	497621	496395	gil143002	proton glutamate symport protein [Bacillus caldotenax]	59	34

Borrelia burgdorferi - Putative coding regions of novel proteins similar to know proteins

1	941	885060	886019	gil1685110	tetrahydrofolate dehydrogenase/cyclohydrolase [Streptococcus thermophilus]	59	36
1	40	50348	48951	gil1574003	pantothenate permease (panF) [Haemophilus influenzae]	58	38
1	76	90160	89534	gil1303791	YqeJ [Bacillus subtilis]	58	32
1	116	115845	115654	gnllPIDle275892	T06E6.f [Caenorhabditis elegans]	58	37
1	179	173515	173009	gil1573163	hypothetical [Haemophilus influenzae]	58	37
1	197	191904	189634	gil1066850	putative [Rhodobacter capsulatus]	58	37
1	229	215111	214563	gil1573441	oxygen-independent coproporphyrinogen III oxidase (hemN) [Haemophilus influenzae]	58	34
1	257	238952	241873	gil1041785	rhoptry protein [Plasmodium yoelii]	58	30
1	440	421010	420792	gil1674178	(AE000047) Mycoplasma pneumoniae, MG246 homolog, from M. genitalium [Mycoplasma pneumoniae]	58	37
1	557	533653	534750	gil974332	NAD(P)H-dependent dihydroxyacetone-phosphate reductase [Bacillus subtilis]	58	41
1	586	557259	559370	gil153062	helicase [Staphylococcus aureus]	58	41
1	623	591542	592435	gil1653618	hypothetical protein [Synechocystis sp.]	58	35
1	728	683208	684104	gil790935	fliG [Treponema denticola]	58	31
1	796	750629	749508	gil1574412	alanine racemase, biosynthetic (alr) [Haemophilus influenzae]	58	29
1	823	778475	778723	gil1209836	minus strand repeat motif-containing gene [Borrelia burgdorferi]	58	22
1	830	786540	788225	gil1574150	ribosomal protein S1 (rpS1) [Haemophilus influenzae]	58	34
1	842	796255	796019	gnllPIDle243474	ORF YGR089w [Saccharomyces cerevisiae]	58	35
1	883	834332	834520	gil1575792	low Mr GTP-binding protein Rab32 [Homo sapiens]	58	43
1	905	853953	854435	gil1303823	YqfG [Bacillus subtilis]	58	34
1	919	863594	862875	gil1256625	putative [Bacillus subtilis]	58	34
1	921	865297	864725	gil1054584	putative protein highly homologous to E. coli RNase HII [Magnetospirillum sp.]	58	42
1	196	189636	187702	gil496484	tlpC gene product [Bacillus subtilis]	57	32
1	262	249142	248192	gil46605	lacC polypeptide (AA 1-310) [Staphylococcus aureus]	57	41
1	311	300776	301660	gil467431	high level kasgamycin resistance [Bacillus subtilis]	57	35
1	365	358725	358495	gil396943	early protein [Human papillomavirus type 19]	57	38
1	386	378249	378025	gil45986	NAD synthetase [Rhodobacter capsulatus]	57	32

Borrelia burgdorferi - Putative coding regions of novel proteins similar to know proteins

1	399	394690	394247	gil1592085	M. jannaschii predicted coding region MJ1437 [Methanococcus jannaschii]	57	31
1	402	397512	396193	gil1732241	GTP-binding protein [Treponema pallidum]	57	36
1	533	504504	505022	gil786163	Ribosomal Protein L10 [Bacillus subtilis]	57	29
1	735	689992	689096	gil1303855	YqgH [Bacillus subtilis]	57	30
1	794	745857	747644	gil11665	ORF2136 [Marchantia polymorpha]	57	33
1	814	768735	771866	gil1573914	acriflavine resistance protein (acrB) [Haemophilus influenzae]	57	31
1	821	776835	778244	gil1591660	histidyl-tRNA synthetase [Methanococcus jannaschii]	57	36
1	834	790907	791752	gil1518661	elongation factor Ts [Chlamydia trachomatis]	57	36
1	836	792328	793038	gil1573941	hypothetical [Haemophilus influenzae]	57	36
1	848	799086	799670	gil537044	50S ribosomal subunit protein L9 [Escherichia coli]	57	39
1	849	799668	801041	gil1001271	replicative DNA helicase [Synechocystis sp.]	57	33
1	851	802510	803742	pirlA64092lA 64092	acetyl coenzyme A acetyltransferase (thiolase) (fadA) homolog - Haemophilus influenzae (strain Rd KW20)	57	43
1	855	805240	806952	gil1499620	M. jannaschii predicted coding region MJ0798 [Methanococcus jannaschii]	57	36
1	922	865347	867809	gil1237015	ORF4 [Bacillus subtilis]	57	38
1	12	17611	14156	pirlB30565lB 30565	phospholipase C (EC 3.1.4.3) precursor - Clostridium bifermentans	56	28
1	26	35530	34277	gil1657594	exonuclease SbcD [Escherichia coli]	56	39
1	59	68915	68271	gil148876	probable com101A gene [Haemophilus influenzae]	56	26
1	79	91821	91480	gil1139633	large tegument protein [Human herpesvirus 7]	56	30
1	112	113768	113571	gnllPIDle2469 33	ORF YPL216w [Saccharomyces cerevisiae]	56	35
1	147	142606	143988	gil642030	NADH oxidase [Serpulina hyodysenteriae]	56	36
1	153	148561	149100	gil1499018	M. jannaschii predicted coding region MJ0240 [Methanococcus jannaschii]	56	29
1	169	165431	164388	gil1573431	aminodeoxychorismate lyase (pabC) [Haemophilus influenzae]	56	41
1	183	176655	175432	gil143841	xylose repressor [Bacillus subtilis]	56	38
1	312	301170	300922	gnllPIDle2202 40	red alga1 chloroplast [Plasmodium falciparum]	56	36
1	317	308362	306992	gil1574691	UDP-N-acetylmuramoylalanine-D-glutamate ligase (murD) [Haemophilus influenzae]	56	40

Borrelia burgdorferi - Putative coding regions of novel proteins similar to know proteins

1	448	426477	426133	gil467410	unknown [Bacillus subtilis]	56	28
1	456	432628	434457	gil142521	deoxyribodipyrimidine photolyase [Bacillus subtilis]	56	34
1	460	438178	437312	gil882453	ORF_f286; alternate name yggB; orf4 of X14436 [Escherichia coli]	56	31
1	469	441309	443438	gil148316	NaH-antiporter protein [Enterococcus hirae]	56	32
1	608	574772	574951	gil1019630	NADH dehydrogenase subunit 2 [Paramecium aurelia]	56	37
1	699	659498	660055	gil1372995	OrfH [Borrelia burgdorferi]	56	24
1	757	713509	713712	gil861327	F31D5.5 gene product [Caenorhabditis elegans]	56	40
1	791	741305	742837	gil1651873	4-alpha-glucanotransferase [Synechocystis sp.]	56	43
1	822	779478	778291	gil1500309	M. jannaschii predicted coding region MJ1428 [Methanococcus jannaschii]	56	28
1	967	907556	908932	gil1749528	similar to Saccharomyces cerevisiae probable UTP-glucose-1-phosphate uridylyltransferase, SWISS-PROT Accession Number P32861 [Schizosaccharomyces pombe]	56	37
1	39	48953	48048	gil1045895	hypothetical protein (SP:P23851) [Mycoplasma genitalium]	55	41
1	131	132989	131967	gil1574007	nitrogen fixation nifR3 protein (nifR3) (PIR:S49971) [Haemophilus influenzae]	55	39
1	152	148506	147148	gil1653100	Na+ -ATPase subunit J [Synechocystis sp.]	55	31
1	359	352690	353313	gil1213334	OrfX; hypothetical 22.5 KD protein downstream of type IV prepilin leader peptidase gene; Method: conceptual translation supplied by author [Vibrio vulnificus]	55	33
1	361	355510	354140	gil882698	L-fuculose kinase [Escherichia coli]	55	44
1	515	488398	487652	gil397486	endonuclease G [Bos taurus]	55	33
1	551	526427	525285	gil558266	orf gene product [Wolinella succinogenes]	55	30
1	570	543745	544482	gil1303811	YqeU [Bacillus subtilis]	55	33
1	579	551201	551494	gil290487	50S ribosomal subunit protein L28 [Escherichia coli]	55	37
1	584	555359	556063	gil1592301	M. jannaschii predicted coding region MJ0687 [Methanococcus jannaschii]	55	32
1	706	665310	665936	gil403984	deoxyguanosine kinase/deoxyadenosine kinase(I) subunit [Lactobacillus acidophilus]	55	38
1	771	722876	723538	gil1736440	O-sialoglycoprotein endopeptidase (EC 3.4.24.57) (Glycoprotease). [Escherichia coli]	55	39
1	786	736537	737187	gil1589778	SPINDLY [Arabidopsis thaliana]	55	34

Borrelia burgdorferi - Putative coding regions of novel proteins similar to know proteins

1	810	765243	766130	gil984805	glycine betaine-binding protein precursor [Bacillus subtilis]	55	35
1	871	823341	823790	gil1590959	ATP synthase, subunit K [Methanococcus jannaschii]	55	34
1	898	847660	849462	gil1517942	aminopeptidase P [Sus scrofa]	55	46
1	924	867811	868236	gil1142660	POM1 [Plasmodium chabaudi chabaudi]	55	41
1	927	870905	870039	gil534839	CheR [Rhizobium meliloti]	55	32
1	964	904091	903900	gil312694	ARS-binding factor 1 [Kluyveromyces marxianus]	55	50
1	33	44068	43124	gil146860	delta-2-isopentenyl pyrophosphate transferase [Escherichia coli]	54	31
1	63	79094	74679	gil415736	Orf635 gene product [Euglena gracilis]	54	37
1	192	184282	182969	gil151259	HMG-CoA reductase (EC 1.1.1.88) [Pseudomonas mevalonii]	54	35
1	200	194105	192951	gil1045800	ribose transport system permease protein [Mycoplasma genitalium]	54	29
1	224	210749	212320	gil1591243	M. jannaschii predicted coding region MJ0539 [Methanococcus jannaschii]	54	45
1	256	237491	238954	gnlIPIDle245024	unknown [Mycobacterium tuberculosis]	54	34
1	260	245698	247542	gil1574782	exodeoxyribonuclease V (recD) [Haemophilus influenzae]	54	36
1	320	311333	312133	gil1209528	D,D-carboxypeptidase [Enterococcus faecalis]	54	40
1	610	577096	579909	gil1499043	M. jannaschii predicted coding region MJ0263 [Methanococcus jannaschii]	54	30
1	765	720685	719999	gil290216	[bride of sevenless] gene product [Drosophila virilis]	54	25
1	789	739607	739996	gil473804	'dosage-dependent dnaK suppressor protein' [Escherichia coli]	54	35
1	845	797932	798366	gil1045767	ribosomal protein S6 [Mycoplasma genitalium]	54	35
1	951	894898	893912	gil1303842	YqfU [Bacillus subtilis]	54	28
1	86	96019	97032	gil405550	flagellar P-ring protein [Pseudomonas putida]	53	40
1	89	98331	99215	gil912478	No definition line found [Escherichia coli]	53	35
1	164	159533	158562	gil1499620	M. jannaschii predicted coding region MJ0798 [Methanococcus jannaschii]	53	39
1	250	234276	232861	gil1303989	YqkI [Bacillus subtilis]	53	28
1	278	266053	267426	gil1749686	similar to Saccharomyces cerevisiae unknown, EMBL Accession Number Z68194 [Schizosaccharomyces pombe]	53	28
1	302	292150	294309	gil1015945	methyl accepting chemotaxis homolog [Treponema denticola]	53	31
1	364	358298	357702	gil1499620	M. jannaschii predicted coding region MJ0798 [Methanococcus jannaschii]	53	41

Borrelia burgdorferi - Putative coding regions of novel proteins similar to know proteins

1	514	486253	486888	gil940842	orf 06111 gene product [Saccharomyces cerevisiae]	53	28
1	567	541832	540684	gil1165254	YlxH [Borrelia burgdorferi]	53	33
1	621	590418	591032	gil1592021	cell division protein J [Methanococcus jannaschii]	53	32
1	805	759748	758537	gil1732203	GlcNAc 6-P deacetylase [Vibrio furnissii]	53	35
1	854	804825	805298	gil1303915	YqhZ [Bacillus subtilis]	53	30
1	884	835705	834944	gil1574399	H. influenzae predicted coding region HI1555 [Haemophilus influenzae]	53	26
1	48	58236	56944	gil1652686	hypothetical protein [Synechocystis sp.]	52	26
1	53	63264	62383	gil42219	P35 gene product (AA 1 - 314) [Escherichia coli]	52	34
1	56	66168	65665	gil1151158	repeat organellar protein [Plasmodium chabaudi]	52	37
1	95	102255	102746	gil1574136	colicin V production protein (pur regulon) (cvpA) [Haemophilus influenzae]	52	24
1	117	115800	116879	gil288998	secA gene product [Antithamnion sp.]	52	34
1	220	208898	208446	gil1652602	hypothetical protein [Synechocystis sp.]	52	25
1	285	274152	272764	gnlPIDle255128	trigger factor [Bacillus subtilis]	52	28
1	352	344946	346532	gil160299	glutamic acid-rich protein [Plasmodium falciparum]	52	26
1	368	361087	361800	gil216861	24K membrane protein [Pseudomonas aeruginosa]	52	32
1	376	368462	367695	gil147213	phnP protein [Escherichia coli]	52	47
1	381	373209	372412	gil467459	unknown [Bacillus subtilis]	52	28
1	437	418141	416768	gil1591425	hypothetical protein (GP:X91006_2) [Methanococcus jannaschii]	52	27
1	439	420801	420166	gil1674178	(AE000047) Mycoplasma pneumoniae, MG246 homolog, from M. genitalium [Mycoplasma pneumoniae]	52	40
1	474	443436	443798	gil1573287	aspartyl-tRNA synthetase (aspS) [Haemophilus influenzae]	52	35
1	583	555235	553802	gil496254	fibronectin/fibrinogen-binding protein [Streptococcus pyogenes]	52	29
1	759	715852	715610	gil397703	dihydroorotate dehydrogenase [Plasmodium falciparum]	52	30
1	797	751384	750674	gil1063419	S2 gene product [Borrelia burgdorferi]	52	27
1	820	776768	774852	gil580936	SpoVD [Bacillus subtilis]	52	33
1	869	820887	821516	gil1592298	ATP synthase, subunit D [Methanococcus jannaschii]	52	29
1	888	839581	838106	gil1151158	repeat organellar protein [Plasmodium chabaudi]	52	30
1	916	862856	862110	gil1256625	putative [Bacillus subtilis]	52	25
1	67	83112	81610	gil587604	beta subunit RNA polymerase [Plasmodium falciparum]	51	29

Borrelia burgdorferi - Putative coding regions of novel proteins similar to know proteins

1	150	147190	146360	gil520844	orf4 [Bacillus subtilis]	51	26
1	194	186516	185275	gil211931	3-hydroxy-3-methylglutaryl-CoA synthase [Gallus gallus]	51	29
1	300	288759	289676	gil142833	ORF2 [Bacillus subtilis]	51	33
1	371	362209	362874	gil1698880	protein antigen LmSTI1 [Leishmania major]	51	27
1	464	438943	439497	gil1591434	chromate resistance protein A [Methanococcus jannaschii]	51	29
1	819	772935	774842	gnllPIDle239057	AMP-binding protein [Brassica napus]	51	29
1	926	869257	869955	gil633996	a negative regulator of pho regulon [Pseudomonas aeruginosa]	51	26
1	45	54760	54062	gil505363	ORF2 [Salmonella typhimurium]	50	32
1	94	101155	102261	gil39995	phospho-N-acetylmuramoyl-pentapeptide- transferase [Bacillus subtilis]	50	29
1	118	118397	117096	gil1762996	RING-finger protein [Helicoverpa armigera nucleopolyhedrovirus]	50	25
1	155	151159	150506	gil893358	PgsA [Bacillus subtilis]	50	35
1	239	224187	224744	gil1303843	YqfV [Bacillus subtilis]	50	29
1	274	265044	264040	gnllPIDle276778	unknown [Mycobacterium tuberculosis]	50	32
1	287	276164	274710	gil147140	peptidase D [Escherichia coli]	50	28
1	310	299525	300778	gil1652202	ComE [Synechocystis sp.]	50	33
1	349	342477	341581	gnllPIDle220245	frameshift [Plasmodium falciparum]	50	30
1	457	435120	434509	gil144839	beta-galactosidase [Thermoanaerobacterium thermosulfurigenes]	50	29
1	479	448691	447948	gil580905	B.subtilis genes rpmH, rnpA, 50kd, gidA and gidB [Bacillus subtilis]	50	32
1	680	640194	641039	gil882579	CG Site No. 29739 [Escherichia coli]	50	31
1	737	690152	690400	gil1086864	T03G11.2 gene product [Caenorhabditis elegans]	50	39
1	752	708130	709662	gil40162	murE gene product [Bacillus subtilis]	50	30
1	360	353288	354157	gil343314	involucrin [Saguinus oedipus]	49	20
1	44	54046	53216	gil505363	ORF2 [Salmonella typhimurium]	48	21
1	122	119896	120774	gil405908	yejE [Escherichia coli]	48	29
1	161	157504	156653	gil143213	putative [Bacillus subtilis]	48	23
1	316	305940	306995	gil1762962	FemA [Staphylococcus simulans]	48	28
1	459	436152	437315	gil1001478	hypothetical protein [Synechocystis sp.]	48	25

Borrelia burgdorferi - Putative coding regions of novel proteins similar to know proteins

1	628	596267	596566	gil156218	putative [Caenorhabditis elegans]	48	32
1	694	655069	654452	gil1574476	dedA protein (dedA) [Haemophilus influenzae]	48	22
1	731	686392	686129	gil915207	gastric mucin [Sus scrofa]	48	27
1	893	844951	843476	gnl|PID|e220245	frameshift [Plasmodium falciparum]	48	32
1	62	74673	72196	gil1766042	outer membrane protein [Neisseria gonorrhoeae]	47	30
1	103	107896	108780	gil1256885	P24A protein (unknown function) (Swiss Prot. accession number P32802) [Saccharomyces cerevisiae]	47	27
1	187	181111	180215	gil1184118	mevalonate kinase [Methanobacterium thermoautotrophicum]	47	30
1	204	195930	196616	gil1573552	phosphoglycolate phosphatase, chromosomal (SP:P40852) [Haemophilus influenzae]	47	21
1	265	251835	251098	gil1209847	repeat motif-containing gene [Borrelia burgdorferi]	47	30
1	334	325837	326529	gil1591893	uridylate kinase [Methanococcus jannaschii]	47	26
1	356	349581	349991	gil849173	Probable essential component of the nucleoskeleton (Swiss Prot. accession number P32380) [Saccharomyces cerevisiae]	47	27
1	490	460559	459834	gil1592264	type I restriction enzyme [Methanococcus jannaschii]	47	34
1	526	499992	499264	gil710551	ankyrin 3 [Mus musculus]	47	29
1	577	549541	548390	gnl|PID|e220245	frameshift [Plasmodium falciparum]	47	27
1	744	701189	699441	gnl|PID|e160436	orfA gene product [Borrelia burgdorferi]	47	23
1	755	713050	710765	pir|S41649|S41649	DNA polymerase - Plasmodium falciparum	47	22
1	761	717229	718392	gil1500309	M. jannaschii predicted coding region MJ1428 [Methanococcus jannaschii]	47	37
1	813	767745	768737	pir|G64100|G64100	membrane fusion protein (mtrC) homolog - Haemophilus influenzae (strain Rd KW20)	47	23
1	824	779587	780546	gil687844	contains TPR domain-like repeats [Caenorhabditis elegans]	47	28
1	881	834283	833015	gil1574393	H. influenzae predicted coding region HI1548 [Haemophilus influenzae]	47	24
1	886	837236	836199	gil887563	serine/threonine-protein kinase [Plasmodium falciparum]	47	30
1	47	57001	55880	gil1652686	hypothetical protein [Synechocystis sp.]	46	23
1	160	156659	156171	gil13261	ORF4 protein (AA 1-156) [Paramecium aurelia]	46	30

Borrelia burgdorferi - Putative coding regions of novel proteins similar to know proteins

1	249	231765	232829	gil1142681	Lpp38 [Pasteurella haemolytica]	46	28
1	329	323695	322838	gil562039	NADH dehydrogenase, subunit 2 [Acanthamoeba castellanii]	46	20
1	336	329090	327303	gil457146	rhoptry protein [Plasmodium yoelii]	46	18
1	442	422511	421747	gil1591598	hypothetical protein (GP:U19364_6) [Methanococcus jannaschii]	46	27
1	452	428632	429375	pirlS41649lS41649	DNA polymerase - Plasmodium falciparum	46	21
1	573	545081	545596	gil1022328	Four tandem repeats of a DNA-binding domain known as the AT-hook are found at the carboxy terminus of CarD. This protein has been purified and found to bind in vitro to a promoter region [Myxococcus xanthus]	46	28
1	617	586903	587865	gnllPIDle33329	ND5 protein [Ascaris suum]	46	29
1	708	668290	666710	gil1573271	apolipoprotein N-acyltransferase (cute) [Haemophilus influenzae]	46	32
1	790	741189	740008	gil458015	TpN50 precursor [Treponema pallidum]	46	29
1	892	843474	841147	gil1574537	outer membrane integrity protein (tolA) [Haemophilus influenzae]	46	19
1	903	853463	852741	gil806562	nebulin [Homo sapiens]	46	26
1	968	908917	909948	gil438455	possible N-terminal signal sequence; mature protein may be membrane-anchored and start at Cys-17. 17.5% identity over 354-aa overlap with Candida pelliculosa beta-glucosidase.; putative [Bacillus subtilis]	46	29
1	208	197467	198516	gil220578	open reading frame [Mus musculus]	45	27
1	462	438197	438949	gil687689	similar to a chromate resistance protein (ChrA) from A. eutrophus, Swiss-Prot Accession Number P17551 [Synechococcus sp.]	45	23
1	742	698657	695295	gil1499043	M. jannaschii predicted coding region MJ0263 [Methanococcus jannaschii]	45	23
1	90	99196	98756	gil303895	ORF 8: This ORF is required for the secretion of IpaB, IpaC and IpaD [Plasmid pMYSH6000]	44	26
1	253	235698	234343	gil143245	Na+/H+ antiporter [Bacillus firmus]	44	26
1	709	668406	670430	gnllPIDle236483	F54G8.4 [Caenorhabditis elegans]	44	31
1	850	802490	801045	gnllPIDle220245	frameshift [Plasmodium falciparum]	44	25
1	458	436119	435118	gil1303799	YqeN [Bacillus subtilis]	43	19

Borrelia burgdorferi - Putative coding regions of novel proteins similar to know proteins

1	859	810560	809967	pir|S17998|S17998	gene COX1 intron 4 protein - yeast (Kluyveromyces marxianus var. lactis) mitochondrion (SGC2)	43	30
1	935	881179	879701	gil1045905	no score generated - score shown is bogus [Mycoplasma genitalium]	43	27
1	319	311250	309877	gil1591425	hypothetical protein (GP:X91006_2) [Methanococcus jannaschii]	42	19
1	618	587863	588672	gil1045801	hypothetical protein (SP:P32720) [Mycoplasma genitalium]	42	24
1	625	593472	594572	gil343962	VAR1 protein [Candida glabrata]	42	25
1	93	100191	101021	gil413976	ipa-52r gene product [Bacillus subtilis]	41	26
1	574	545523	546581	gnl|PID|e1632 6	MURF2 protein (AA 1-348) [Crithidia fasciculata]	41	26
1	740	693458	692403	gil1151158	repeat organellar protein [Plasmodium chabaudi]	41	20
1	3	5792	6796	gil1256888	Similar to chromosome segregation protein Smc1p of S. cerevisiae (GenBank accession number L00602), chromosome segregation protein Cut3p of S. pombe (Swiss Prot. accession number P41004), and C. elegans hypothetical proteins R13G10.1 (GenBank	40	24
1	228	214440	214742	gil1150836	neural specific DNA binding protein [Xenopus laevis]	40	28
1	318	309735	308377	gil1591425	hypothetical protein (GP:X91006_2) [Methanococcus jannaschii]	40	19
1	453	431037	429700	gil499647	[Mus musculus (strain C3HF/RL) ORF mRNA, complete cds.], gene product [Mus musculus]	40	24
1	795	747813	749516	gil304179	wall-associated protein [Bacillus subtilis]	40	35
1	966	907336	905528	gil1151158	repeat organellar protein [Plasmodium chabaudi]	40	26

TABLE 2. Borrelia burgdorferi - Coding regions containing know proteins

Contig ID	Orf ID	Start (nt)	Stop (nt)	match acession	match gene name	percent ident	HSP nt length
1	69	86349	85018	gb|L32144|	Borrelia burgdorferi peptidyl-tRNA hydrolase-like protein (pth) gene homologue, complete cds	100	220
1	70	86918	86340	gb|L32144|	Borrelia burgdorferi peptidyl-tRNA hydrolase-like protein (pth) gene homologue, complete cds	100	579
1	71	87573	86911	gb|L32144|	Borrelia burgdorferi peptidyl-tRNA hydrolase-like protein (pth) gene homologue, complete cds	100	129
1	124	123885	121759	gb|M60802|	B.burgdorferei immunogen gene, 5' flank	99	2127
1	126	127421	125700	emb|X91965| BBATPBP	B.burgdorferi abp gene	97	284
1	137	136332	139151	gb|L31424|	Borrelia burgdorferi (clone BbK3.11) phoA fusion protein gene, partial cds	98	248
1	138	138676	138515	gb|L31424|	Borrelia burgdorferi (clone BbK3.11) phoA fusion protein gene, partial cds	96	60
1	165	160705	159932	gb|U17591|	Borrelia burgdorferi primary sigma factor (rpoD) gene, complete cds	100	774
1	166	162604	160703	gb|U17591|	Borrelia burgdorferi primary sigma factor (rpoD) gene, complete cds	100	1902
1	167	162835	162602	gb|U17591|	Borrelia burgdorferi primary sigma factor (rpoD) gene, complete cds	99	232
1	168	164397	162811	gb|U17591|	Borrelia burgdorferi primary sigma factor (rpoD) gene, complete cds	99	1216
1	210	198495	199028	gb|U61498|	Borrelia burgdorferi CheA (cheA) gene, partial cds, CheW (cheW), CheX (cheX) and CheY (cheY) genes, complete cds	98	127
1	211	199527	199069	gb|U61498|	Borrelia burgdorferi CheA (cheA) gene, partial cds, CheW (cheW), CheX (cheX) and CheY (cheY) genes, complete cds	99	459
1	212	200067	199549	gb|U61498|	Borrelia burgdorferi CheA (cheA) gene, partial cds, CheW (cheW), CheX (cheX) and CheY (cheY) genes, complete cds	99	519
1	213	201455	200046	gb|U61498|	Borrelia burgdorferi CheA (cheA) gene, partial cds, CheW (cheW), CheX (cheX) and CheY	99	1410

					(cheY) genes, complete cds		
1	214	204053	201453	gb|U28962|	Borrelia burgdorferi histidine kinase (cheA) gene, complete cds	99	2601
1	215	205173	204115	gb|U62900|	Borrelia burgdorferi flagellar filament outsheath protein (flaA) gene, complete cds, and chemotaxis histidine kinase (cheA) gene, partial cds	98	1059
1	216	205780	205220	gb|U62900|	Borrelia burgdorferi flagellar filament outsheath protein (flaA) gene, complete cds, and chemotaxis histidine kinase (cheA) gene, partial cds	98	277
1	235	220591	220232	emb|X65139| BBHSP60	B.burgdorferi hsp60 gene for 60kDa heat shock protein	98	139
1	236	222288	220594	emb|X65139| BBHSP60	B.burgdorferi hsp60 gene for 60kDa heat shock protein	99	1695
1	237	222396	224114	emb|X54059| BBGROEL	Borrelia burgdorferi groEL gene for a common antigen	94	416
1	241	227310	225754	gb|L31417|	Borrelia burgdorferi (clone BbK1.4) phoA fusion protein gene, partial cds	99	231
1	295	281749	283629	emb|X87725| BBDNA66KD	B.burgdorferi p66 gene for 66kDa protein	100	1881
1	296	284438	283683	gb|M58431|	Borrelia burgdorferei PCR target sequence	93	356
1	315	303735	305942	gb|L32146|	Borrelia burgdorferi methionyl tRNA synthetase (metG) gene, partial cds	99	345
1	337	330246	329077	gb|U60236|	Borrelia burgdorferi response regulator gene, partial cds	100	191
1	338	332455	330299	gb|L39965|	Borrelia burgdorferi histidine kinase (cheA) gene, complete cds	99	1361
1	344	336894	335473	gb|U51878|	Borrelia burgdorferi phosphotransferase enzyme II (crr) gene, hsp90 (hptg) gene, complete cds	100	95
1	345	336851	338830	gb|U51878|	Borrelia burgdorferi phosphotransferase enzyme II (crr) gene, hsp90 (hptg) gene, complete cds	99	1980
1	346	339458	338868	gb|U51878|	Borrelia burgdorferi phosphotransferase enzyme	100	591

Borrelia burgdorferi - Coding regions containing know proteins

					II (crr) gene, hsp90 (hptg) gene, complete cds		
1	388	378955	379590	gb|M96847|	Borrelia burgdorferi GrpE protein homologue gene, DnaK protein homologue gene, and DnaJ protein homologue gene, complete cds's	100	636
1	389	379566	381521	emb|X67646| BBHSPRO	B.burgdorferi dnaK gene for heat-shock protein	100	1956
1	390	381512	381943	gb|M97914|	Borrelia burgdorferi DnaJ gene, complete cds	97	424
1	391	381907	382617	gb|M96847|	Borrelia burgdorferi GrpE protein homologue gene, DnaK protein homologue gene, and DnaJ protein homologue gene, complete cds's	100	687
1	392	382656	383360	gb|M96847|	Borrelia burgdorferi GrpE protein homologue gene, DnaK protein homologue gene, and DnaJ protein homologue gene, complete cds's	95	144
1	393	383005	382688	gb|M96847|	Borrelia burgdorferi GrpE protein homologue gene, DnaK protein homologue gene, and DnaJ protein homologue gene, complete cds's	95	144
1	394	384408	383416	gb|U82978|	Borrelia burgdorferi phenylalanyl-tRNA synthetase alpha subunit (pheS), phenylalanyl-tRNA synthetase beta subunit (pheT) and thioredoxin reductase (trxB) genes, complete cds	99	956
1	395	384799	384467	gb|U82978|	Borrelia burgdorferi phenylalanyl-tRNA synthetase alpha subunit (pheS), phenylalanyl-tRNA synthetase beta subunit (pheT) and thioredoxin reductase (trxB) genes, complete cds	99	292
1	396	386169	384733	gb|U82978|	Borrelia burgdorferi phenylalanyl-tRNA synthetase alpha subunit (pheS), phenylalanyl-tRNA synthetase beta subunit (pheT) and thioredoxin reductase (trxB) genes, complete cds	99	1416
1	397	387733	386144	gb|U82978|	Borrelia burgdorferi phenylalanyl-tRNA synthetase alpha subunit (pheS), phenylalanyl-tRNA synthetase beta subunit (pheT) and thioredoxin reductase (trxB) genes, complete cds	99	1220
1	398	394257	387727	gb|U82978|	Borrelia burgdorferi phenylalanyl-tRNA synthetase alpha subunit (pheS), phenylalanyl-	99	230

Borrelia burgdorferi - Coding regions containing know proteins

					tRNA synthetase beta subunit (pheT) and thioredoxin reductase (trxB) genes, complete cds		
1	421	408559	407981	gb|M28681|	B.burgdorferei promoter region DNA	98	152
1	427	411019	410132	gb|U78193|	Borrelia burgdorferi tuf-s10 operon: elongation factor (tuf), ribosomal proteins S10 (rpsJ), L3 (rplC), L4 (rplD), L23 (rplW), L2 (rplB), S19 (rpsS), and L22 (rplV)genes, complete cds, and S3 (rpsC) gene, partial cds	99	287
1	428	411388	411017	gb|U78193|	Borrelia burgdorferi tuf-s10 operon: elongation factor (tuf), ribosomal proteins S10 (rpsJ), L3 (rplC), L4 (rplD), L23 (rplW), L2 (rplB), S19 (rpsS), and L22 (rplV)genes, complete cds, and S3 (rpsC) gene, partial cds	96	357
1	429	411676	411386	gb|U78193|	Borrelia burgdorferi tuf-s10 operon: elongation factor (tuf), ribosomal proteins S10 (rpsJ), L3 (rplC), L4 (rplD), L23 (rplW), L2 (rplB), S19 (rpsS), and L22 (rplV)genes, complete cds, and S3 (rpsC) gene, partial cds	99	291
1	430	412531	411674	gb|U78193|	Borrelia burgdorferi tuf-s10 operon: elongation factor (tuf), ribosomal proteins S10 (rpsJ), L3 (rplC), L4 (rplD), L23 (rplW), L2 (rplB), S19 (rpsS), and L22 (rplV)genes, complete cds, and S3 (rpsC) gene, partial cds	98	858
1	431	412852	412529	gb|U78193|	Borrelia burgdorferi tuf-s10 operon: elongation factor (tuf), ribosomal proteins S10 (rpsJ), L3 (rplC), L4 (rplD), L23 (rplW), L2 (rplB), S19 (rpsS), and L22 (rplV)genes, complete cds, and S3 (rpsC) gene, partial cds	99	324
1	432	413487	412846	gb|U78193|	Borrelia burgdorferi tuf-s10 operon: elongation factor (tuf), ribosomal proteins S10 (rpsJ), L3 (rplC), L4 (rplD), L23 (rplW), L2 (rplB), S19 (rpsS), and L22 (rplV)genes, complete cds, and S3 (rpsC) gene, partial cds	99	642
1	433	414117	413485	gb|U78193|	Borrelia burgdorferi tuf-s10 operon: elongation factor (tuf), ribosomal proteins S10 (rpsJ), L3	99	633

					(rplC), L4 (rplD), L23 (rplW), L2 (rplB), S19 (rpsS), and L22 (rplV)genes, complete cds, and S3 (rpsC) gene, partial cds		
1	434	414464	414141	gb|U78193|	Borrelia burgdorferi tuf-s10 operon: elongation factor (tuf), ribosomal proteins S10 (rpsJ), L3 (rplC), L4 (rplD), L23 (rplW), L2 (rplB), S19 (rpsS), and L22 (rplV)genes, complete cds, and S3 (rpsC) gene, partial cds	100	324
1	435	415714	414503	gb|L23125|	Borrelia burgdorferi elongation factor EF-Tu (tuf) gene, complete cds	100	1212
1	481	450681	450310	gb|U04527|	Borrelia burgdorferi 212 DNA gyrase b subunit (gyrB) and ribonuclease P protein component (rnpA) genes, partial cds, DnaA protein (dnaA), DNA polymerase III beta subunit (dnaN), and ribosomal protein L34 (rpmH) genes, complete cds	100	148
1	482	450820	450650	gb|U04527|	Borrelia burgdorferi 212 DNA gyrase b subunit (gyrB) and ribonuclease P protein component (rnpA) genes, partial cds, DnaA protein (dnaA), DNA polymerase III beta subunit (dnaN), and ribosomal protein L34 (rpmH) genes, complete cds	100	171
1	483	451208	450897	gb|U04527|	Borrelia burgdorferi 212 DNA gyrase b subunit (gyrB) and ribonuclease P protein component (rnpA) genes, partial cds, DnaA protein (dnaA), DNA polymerase III beta subunit (dnaN), and ribosomal protein L34 (rpmH) genes, complete cds	100	312
1	484	451288	451467	gb|U04527|	Borrelia burgdorferi 212 DNA gyrase b subunit (gyrB) and ribonuclease P protein component (rnpA) genes, partial cds, DnaA protein (dnaA), DNA polymerase III beta subunit (dnaN), and ribosomal protein L34 (rpmH) genes, complete cds	100	180
1	485	452456	451287	gb|U04527|	Borrelia burgdorferi 212 DNA gyrase b subunit	99	1170

					(gyrB) and ribonuclease P protein component (rnpA) genes, partial cds, DnaA protein (dnaA), DNA polymerase III beta subunit (dnaN), and ribosomal protein L34 (rpmH) genes, complete cds		
1	486	454181	452685	gb|U04527|	Borrelia burgdorferi 212 DNA gyrase b subunit (gyrB) and ribonuclease P protein component (rnpA) genes, partial cds, DnaA protein (dnaA), DNA polymerase III beta subunit (dnaN), and ribosomal protein L34 (rpmH) genes, complete cds	100	1497
1	487	454315	456237	gb|U04527|	Borrelia burgdorferi 212 DNA gyrase b subunit (gyrB) and ribonuclease P protein component (rnpA) genes, partial cds, DnaA protein (dnaA), DNA polymerase III beta subunit (dnaN), and ribosomal protein L34 (rpmH) genes, complete cds	98	904
1	488	456228	458681	emb|Z12165|BBGYRAG	B.burgdorferi gyrA gene encoding DNA gyrase subunit A (partial)	96	289
1	496	463825	464394	gb|U03396|	Borrelia burgdorferi B31 Ala-tRNA (alaT), Ile-tRNA (ileT), 16S rRNA, 23S rRNA (rrlA and rrlB), and 5S rRNA (rrfA and rrfB) genes, complete sequence	100	570
1	497	466650	466958	gb|U03396|	Borrelia burgdorferi B31 Ala-tRNA (alaT), Ile-tRNA (ileT), 16S rRNA, 23S rRNA (rrlA and rrlB), and 5S rRNA (rrfA and rrfB) genes, complete sequence	96	210
1	498	467437	468033	gb|U03396|	Borrelia burgdorferi B31 Ala-tRNA (alaT), Ile-tRNA (ileT), 16S rRNA, 23S rRNA (rrlA and rrlB), and 5S rRNA (rrfA and rrfB) genes, complete sequence	99	209
1	499	468167	468433	gb|U03396|	Borrelia burgdorferi B31 Ala-tRNA (alaT), Ile-tRNA (ileT), 16S rRNA, 23S rRNA (rrlA and rrlB), and 5S rRNA (rrfA and rrfB) genes, complete sequence	98	267

1	500	468391	468999	gb|U03396|	Borrelia burgdorferi B31 Ala-tRNA (alaT), Ile-tRNA (ileT), 16S rRNA, 23S rRNA (rrlA and rrlB), and 5S rRNA (rrfA and rrfB) genes, complete sequence	95	386
1	501	470714	470445	gb|M88330|	Borrelia burgdorferi 23S ribosomal RNA gene	100	270
1	502	475597	480090	gb|U03396|	Borrelia burgdorferi B31 Ala-tRNA (alaT), Ile-tRNA (ileT), 16S rRNA, 23S rRNA (rrlA and rrlB), and 5S rRNA (rrfA and rrfB) genes, complete sequence	97	131
1	535	505532	509017	gb|L48488|	Borrelia burgdorferi RNA polymerase beta subunit (rpoB) gene, complete cds, RNA polymerase beta' subunit (rpoC) gene, 5' end of cds	98	2490
1	536	509015	513166	gb|L48488|	Borrelia burgdorferi RNA polymerase beta subunit (rpoB) gene, complete cds, RNA polymerase beta' subunit (rpoC) gene, 5' end of cds	97	76
1	538	513606	514106	gb|U35450|	Borrelia burgdorferi membrane protein D (bmpD) gene, complete cds	100	82
1	539	514120	515229	gb|U35450|	Borrelia burgdorferi membrane protein D (bmpD) gene, complete cds	99	1110
1	540	515472	516605	gb|U49938|	Borrelia burgdorferi potential virulence gene cluster membrane proteins BmpC (bmpC) and BmpA (bmpA), BmpB protein (bmpB), putative protein 4, Mg ion transporter MgtE (mgtE), protein kinase C1 inhibitor PKCI (pkci) genes, complete cds	99	1134
1	541	516641	517666	gb|L24194|	Borrelia burgdorferi immunodominant antigen P39 gene, complete cds	99	1026
1	542	517732	518256	gb|L35050|	Borrelia burgdorferi (clone pB46) membrane lipoprotein A (bmpA) gene, 3' end, membrane lipoprotein (bmpB) gene, 5' end	98	457
1	543	518168	518779	gb|L24194|	Borrelia burgdorferi immunodominant antigen P39 gene, complete cds	99	606

Borrelia burgdorferi - Coding regions containing know proteins

1	544	518856	520316	gb|U49938|	Borrelia burgdorferi potential virulence gene cluster membrane proteins BmpC (bmpC) and BmpA (bmpA), BmpB protein (bmpB), putative protein 4, Mg ion transporter MgtE (mgtE), protein kinase C1 inhibitor PKCI (pkci) genes, complete cds	99	1461
1	545	520349	521734	gb|U49938|	Borrelia burgdorferi potential virulence gene cluster membrane proteins BmpC (bmpC) and BmpA (bmpA), BmpB protein (bmpB), putative protein 4, Mg ion transporter MgtE (mgtE), protein kinase C1 inhibitor PKCI (pkci) genes, complete cds	99	1386
1	546	521752	522204	gb|U49938|	Borrelia burgdorferi potential virulence gene cluster membrane proteins BmpC (bmpC) and BmpA (bmpA), BmpB protein (bmpB), putative protein 4, Mg ion transporter MgtE (mgtE), protein kinase C1 inhibitor PKCI (pkci) genes, complete cds	100	453
1	547	522168	522893	gb|U49938|	Borrelia burgdorferi potential virulence gene cluster membrane proteins BmpC (bmpC) and BmpA (bmpA), BmpB protein (bmpB), putative protein 4, Mg ion transporter MgtE (mgtE), protein kinase C1 inhibitor PKCI (pkci) genes, complete cds	98	130
1	559	535086	534772	emb|X78708| BBYSC1	B.bergdorferi (ZS7) YSC1-like gene	99	314
1	560	536461	535058	emb|X78708| BBYSC1	B.bergdorferi (ZS7) YSC1-like gene	100	1404
1	561	536545	537144	emb|X70826| BBLA7	B.burgdorferi gene for lipoprotein	100	600
1	562	537652	537191	emb|X70826| BBLA7	B.burgdorferi gene for lipoprotein	100	57
1	563	539695	537665	gb|M90084|	Borrelia burgdorferi 22 kD antigen	100	786
1	564	537705	537968	gb|M90084|	Borrelia burgdorferi 22 kD antigen	100	264

1	565	538395	538757	gb|M90084|	Borrelia burgdorferi 22 kD antigen	100	56
1	606	574092	572497	gb|U29143|	Borrelia burgdorferi periplasmic substrate-binding protein homolog (p30) gene, complete cds	92	805
1	607	575817	574204	gb|U29143|	Borrelia burgdorferi periplasmic substrate-binding protein homolog (p30) gene, complete cds	100	84
1	616	585458	586936	gb|L31422|	Borrelia burgdorferi (clone Bb2.13) phoA fusion protein gene, partial cds	100	354
1	629	596586	597983	gb|U43739|	Borrelia burgdorferi fesmid clone 31, complete sequence	100	1185
1	630	597967	599052	emb|X96685| BBCDG	B.burgdorferi cell division genes	99	912
1	631	599050	600153	emb|X96685| BBCDG	B.burgdorferi cell division genes	99	1104
1	632	600183	600932	emb|X96433| BBFTSWQA	B.burgdorferi ftsW, ftsQ & ftsA genes	99	750
1	633	600905	602173	gb|U43739|	Borrelia burgdorferi fesmid clone 31, complete sequence	100	1269
1	634	602171	603394	gb|U43739|	Borrelia burgdorferi fesmid clone 31, complete sequence	100	1224
1	635	603392	604087	gb|U43739|	Borrelia burgdorferi fesmid clone 31, complete sequence	100	696
1	636	604085	605041	gb|L76303|	Borrelia burgdorferi ftsA gene, 3' end of cds, ftsZ, orf230, smf, hslVU, flgBCE, fliEFGHI, flbABC genes, complete cds	98	712
1	637	605039	605599	gb|U43739|	Borrelia burgdorferi fesmid clone 31, complete sequence	100	561
1	638	605535	606938	emb|X96685| BBCDG	B.burgdorferi cell division genes	97	1404
1	639	606936	607379	gb|U43739|	Borrelia burgdorferi fesmid clone 31, complete sequence	100	444
1	640	607382	607861	gb|U43739|	Borrelia burgdorferi fesmid clone 31, complete sequence	100	480

Borrelia burgdorferi - Coding regions containing know proteins

1	641	607831	608208	gb|L76303|	Borrelia burgdorferi ftsA gene, 3' end of cds, ftsZ, orf230, smf, hslVU, flgBCE, fliEFGHI, flbABC genes, complete cds	100	378
1	642	608163	609932	gb|L76303|	Borrelia burgdorferi ftsA gene, 3' end of cds, ftsZ, orf230, smf, hslVU, flgBCE, fliEFGHI, flbABC genes, complete cds	100	1770
1	643	609930	610982	gb|U43739|	Borrelia burgdorferi fesmid clone 31, complete sequence	100	1053
1	644	610961	611917	gb|U43739|	Borrelia burgdorferi fesmid clone 31, complete sequence	100	957
1	645	611915	613246	gb|L76303|	Borrelia burgdorferi ftsA gene, 3' end of cds, ftsZ, orf230, smf, hslVU, flgBCE, fliEFGHI, flbABC genes, complete cds	99	1332
1	646	613222	613674	gb|L76303|	Borrelia burgdorferi ftsA gene, 3' end of cds, ftsZ, orf230, smf, hslVU, flgBCE, fliEFGHI, flbABC genes, complete cds	99	453
1	647	613655	614284	gb|U43739|	Borrelia burgdorferi fesmid clone 31, complete sequence	100	630
1	648	614250	615470	gb|L76303|	Borrelia burgdorferi ftsA gene, 3' end of cds, ftsZ, orf230, smf, hslVU, flgBCE, fliEFGHI, flbABC genes, complete cds	99	1221
1	649	615481	615927	gb|L76303|	Borrelia burgdorferi ftsA gene, 3' end of cds, ftsZ, orf230, smf, hslVU, flgBCE, fliEFGHI, flbABC genes, complete cds	100	447
1	650	615911	617260	gb|U43739|	Borrelia burgdorferi fesmid clone 31, complete sequence	100	1350
1	651	617277	617507	gb|U43739|	Borrelia burgdorferi fesmid clone 31, complete sequence	100	231
1	652	617498	618286	gb|U43739|	Borrelia burgdorferi fesmid clone 31, complete sequence	100	789
1	653	618280	619068	gb|U43739|	Borrelia burgdorferi fesmid clone 31, complete sequence	100	789
1	654	619066	619653	gb|U43739|	Borrelia burgdorferi fesmid clone 31, complete sequence	100	588

1	655	619688	620749	gb|U43739|	Borrelia burgdorferi fesmid clone 31, complete sequence	100	1062
1	656	620789	621136	gb|U43739|	Borrelia burgdorferi fesmid clone 31, complete sequence	100	348
1	657	621114	621755	gb|U43739|	Borrelia burgdorferi fesmid clone 31, complete sequence	100	642
1	658	621742	622530	gb|L75945|	Borrelia burgdorferi flagellar hook protein (flgE), flbD, flagellar motor apparatus (motAB), fliL, fliM, fliZ, flagellar export apparatus (fliPQR, flhB), flhF, flbE genes	99	789
1	659	622028	621822	gb|L75945|	Borrelia burgdorferi flagellar hook protein (flgE), flbD, flagellar motor apparatus (motAB), fliL, fliM, fliZ, flagellar export apparatus (fliPQR, flhB), flhF, flbE genes	100	207
1	660	622515	622802	gb|L75945|	Borrelia burgdorferi flagellar hook protein (flgE), flbD, flagellar motor apparatus (motAB), fliL, fliM, fliZ, flagellar export apparatus (fliPQR, flhB), flhF, flbE genes	100	288
1	661	622811	623623	gb|L75945|	Borrelia burgdorferi flagellar hook protein (flgE), flbD, flagellar motor apparatus (motAB), fliL, fliM, fliZ, flagellar export apparatus (fliPQR, flhB), flhF, flbE genes	99	813
1	662	623007	622819	gb|L75945|	Borrelia burgdorferi flagellar hook protein (flgE), flbD, flagellar motor apparatus (motAB), fliL, fliM, fliZ, flagellar export apparatus (fliPQR, flhB), flhF, flbE genes	100	189
1	663	623706	623458	gb|L75945|	Borrelia burgdorferi flagellar hook protein (flgE), flbD, flagellar motor apparatus (motAB), fliL, fliM, fliZ, flagellar export apparatus (fliPQR, flhB), flhF, flbE genes	100	249
1	664	623608	624741	gb|L75945|	Borrelia burgdorferi flagellar hook protein (flgE), flbD, flagellar motor apparatus (motAB), fliL, fliM, fliZ, flagellar export apparatus (fliPQR, flhB), flhF, flbE genes	99	1134

1	665	624735	626843	gb|L75945|	Borrelia burgdorferi flagellar hook protein (flgE), flbD, flagellar motor apparatus (motAB), fliL, fliM, fliZ, flagellar export apparatus (fliPQR, flhB), flhF, flbE genes	100	2109
1	666	626841	628013	gb|U43739|	Borrelia burgdorferi fesmid clone 31, complete sequence	100	1173
1	667	627998	628912	gb|U43739|	Borrelia burgdorferi fesmid clone 31, complete sequence	99	816
1	668	629151	628807	gb|U43739|	Borrelia burgdorferi fesmid clone 31, complete sequence	100	345
1	669	628910	629398	gb|U43739|	Borrelia burgdorferi fesmid clone 31, complete sequence	100	489
1	670	629371	631305	gb|U43739|	Borrelia burgdorferi fesmid clone 31, complete sequence	100	1935
1	671	631314	631634	gb|U43739|	Borrelia burgdorferi fesmid clone 31, complete sequence	100	286
1	676	636891	635476	gb|M28682|	B.burgdorferei promoter element DNA	100	78
1	687	646982	649420	gb|L77216|	Borrelia burgdorferi (strain B31) protease (lon) gene, complete cds	99	2439
1	688	651760	649409	gb|L77216|	Borrelia burgdorferi (strain B31) protease (lon) gene, complete cds	100	274
1	711	671567	672412	gb|U35673|	Borrelia burgdorferi OrfR gene, partial cds, and S20, Hbb, OrfH and Rho genes, complete cds	99	542
1	712	672418	672744	gb|U35673|	Borrelia burgdorferi OrfR gene, partial cds, and S20, Hbb, OrfH and Rho genes, complete cds	100	327
1	713	672751	673083	gb|U48651|	Borrelia burgdorferi P1G histone-like protein HBbu (hbb) gene, complete cds	100	327
1	714	673081	673491	gb|U35673|	Borrelia burgdorferi OrfR gene, partial cds, and S20, Hbb, OrfH and Rho genes, complete cds	99	411
1	715	673553	675118	gb|U35673|	Borrelia burgdorferi OrfR gene, partial cds, and S20, Hbb, OrfH and Rho genes, complete cds	99	1566
1	716	675164	675424	gb|U35673|	Borrelia burgdorferi OrfR gene, partial cds, and S20, Hbb, OrfH and Rho genes, complete cds	100	106
1	773	724171	723770	gb|U62901|	Borrelia burgdorferi thdF gene, partial cds,	97	67

					putative motility protein (flbF), flagellar hook associated proteins FlgK (flgK) and FlgL (flgL) genes, complete cds		
1	774	723891	724181	gb|U62901|	Borrelia burgdorferi thdF gene, partial cds, putative motility protein (flbF), flagellar hook associated proteins FlgK (flgK) and FlgL (flgL) genes, complete cds	97	77
1	775	725456	724164	gb|U62901|	Borrelia burgdorferi thdF gene, partial cds, putative motility protein (flbF), flagellar hook associated proteins FlgK (flgK) and FlgL (flgL) genes, complete cds	99	780
1	776	727348	725441	gb|U62901|	Borrelia burgdorferi thdF gene, partial cds, putative motility protein (flbF), flagellar hook associated proteins FlgK (flgK) and FlgL (flgL) genes, complete cds	99	1841
1	777	727854	727336	gb|U62901|	Borrelia burgdorferi thdF gene, partial cds, putative motility protein (flbF), flagellar hook associated proteins FlgK (flgK) and FlgL (flgL) genes, complete cds	99	519
1	778	727908	729308	emb|X95669|BBTHDFGID	B.burgdorferi thdF and gidA genes	98	1185
1	779	729284	731176	emb|Z12160|BBGIDAG	B.burgdorferi thdF, gidA and gidB genes	99	1893
1	780	731149	731799	emb|X95668|BBGIDMOXR	B.burgdorferi gidA, gidB and moxR genes	98	381
1	781	731772	732848	emb|X96434|BBGIDBMOX	B.burgdorferi gidB moxR genes and ORF	99	789
1	782	732815	733738	emb|X96434|BBGIDBMOX	B.burgdorferi gidB moxR genes and ORF	100	84
1	798	752154	751372	gb|U51878|	Borrelia burgdorferi phosphotransferase enzyme II (crr) gene, hsp90 (hptg) gene, complete cds	100	57

Borrelia burgdorferi - Coding regions containing know proteins

1	800	754266	753118	gb|AF003354|	Borrelia burgdorferi SecA (secA) gene, complete cds	97	67
1	801	753992	754243	gb|AF003354|	Borrelia burgdorferi SecA (secA) gene, complete cds	96	50
1	802	754283	757015	gb|AF003354|	Borrelia burgdorferi SecA (secA) gene, complete cds	99	2041
1	803	756991	757641	gb|AF003354|	Borrelia burgdorferi SecA (secA) gene, complete cds	100	158
1	806	759909	761930	gb|U66699|	Borrelia burgdorferi flagellar filament cap (filD) gene, complete cds and flagellin protein (flaB) gene, partial cds	98	1149
1	807	762051	763067	emb|X16833| BBFAA	Borrelia burgdorferi gene for flagellum-associated 41kD antigen (flagellin)	99	1017
1	808	763194	764339	emb|X63898| BBHYPP	B.burgdorferi DNA for hypothetical protein	99	1146
1	809	764337	765245	emb|X63898| BBHYPP	B.burgdorferi DNA for hypothetical protein	92	253
1	826	783276	784400	gb|U23457|	Borrelia burgdorferi RecA (recA) gene, complete cds	99	1122
1	827	784412	785182	gb|U23457|	Borrelia burgdorferi RecA (recA) gene, complete cds	82	476
1	828	785142	785918	gb|U23457|	Borrelia burgdorferi RecA (recA) gene, complete cds	99	139
1	907	855179	857182	gb|U28760|	Borrelia burgdorferi glyceraldehyde-3-phosphate dehydrogenase (GAPDH), phosphoglycerate kinase (PGK), triosephosphate isomerase (TPI) genes, complete cds	94	75
1	908	857228	858262	gb|U28760|	Borrelia burgdorferi glyceraldehyde-3-phosphate dehydrogenase (GAPDH), phosphoglycerate kinase (PGK), triosephosphate isomerase (TPI) genes, complete cds	99	1035
1	909	858270	859463	gb|U28760|	Borrelia burgdorferi glyceraldehyde-3-phosphate dehydrogenase (GAPDH), phosphoglycerate kinase (PGK), triosephosphate isomerase (TPI)	99	1194

Borrelia burgdorferi - Coding regions containing know proteins

					genes, complete cds		
1	910	859315	860226	gb|U28760|	Borrelia burgdorferi glyceraldehyde-3-phosphate dehydrogenase (GAPDH), phosphoglycerate kinase (PGK), triosephosphate isomerase (TPI) genes, complete cds	99	912
1	911	860224	860604	gb|U57683|	Borrelia burgdorferi sequence 3' to the triosephosphate isomerase (TPI) gene	97	183
1	912	860645	860316	gb|U57683|	Borrelia burgdorferi sequence 3' to the triosephosphate isomerase (TPI) gene	95	94
1	913	861447	860704	gb|U57684|	Borrelia burgdorferi uracil DNA glycosylase (UDG) gene, partial cds	92	294
1	914	861020	861397	gb|U57684|	Borrelia burgdorferi uracil DNA glycosylase (UDG) gene, partial cds	93	244
1	915	861439	862113	gb|U57684|	Borrelia burgdorferi uracil DNA glycosylase (UDG) gene, partial cds	96	128
1	930	874089	874859	gb|L32861|	Borrelia burgdorferi 1-acyl-sn-glycerol-3-phosphate acetyltransferase (plsC) gene, 3' end; topoisomerase IV beta-subunit (parE) gene, 5' end	99	408
1	931	874877	876679	gb|L32861|	Borrelia burgdorferi 1-acyl-sn-glycerol-3-phosphate acetyltransferase (plsC) gene, 3' end; topoisomerase IV beta-subunit (parE) gene, 5' end	100	252
1	943	887900	886758	emb|Y08885| BBRUVABHL	B.burgdorferi ruvA, ruvB and queA genes	98	293
1	944	887965	888570	emb|Y08885| BBRUVABHL	B.burgdorferi ruvA, ruvB and queA genes	99	606
1	945	888603	889658	emb|Y08885| BBRUVABHL	B.burgdorferi ruvA, ruvB and queA genes	99	1056
1	946	889615	890271	emb|Y08885| BBRUVABH	B.burgdorferi ruvA, ruvB and queA genes	97	342

Borrelia burgdorferi - Coding regions containing know proteins

				L			
1	948	890719	892404	emblY09140l BBPFPB	B.burgdorferi pfpB gene	99	1320
1	950	892893	893909	emblY09142l BBYFII	B.burgdorferi yfiI gene	97	919
1	952	894973	895371	emblX97449l BBPRIAUDK	B.burgdorferi priA and udk genes	88	324
1	953	895308	895991	emblX97449l BBPRIAUDK	B.burgdorferi priA and udk genes	99	684
1	954	897976	895988	emblX97449l BBPRIAUDK	B.burgdorferi priA and udk genes	99	1989
1	955	898577	897963	emblX97449l BBPRIAUDK	B.burgdorferi priA and udk genes	100	152
1	956	899298	898555	emblY09141l BBTRUA	B.burgdorferi truA gene	99	741

TABLE 3.
Borrelia burgdorferi - Putative coding regions of novel proteins not similar to know proteins

Contig ID	ORF ID	Start (nt)	Stop (nt)
1	1	2330	1134
1	2	3317	2934
1	8	11375	13021
1	9	11673	11386
1	10	12925	13629
1	11	13538	14146
1	17	25212	24700
1	18	25782	25357
1	19	26115	25870
1	21	27308	27051
1	22	29628	30458
1	29	40696	41217
1	30	41201	41992
1	31	42542	41985
1	32	42593	42982
1	34	44234	44031
1	38	48041	47079
1	41	49318	49617
1	43	53234	51810
1	50	59737	58208
1	58	68227	67733
1	65	79757	80404
1	66	81516	80401
1	75	89552	88353
1	82	93338	92766
1	85	95207	95854
1	104	108788	108621
1	105	109764	108943
1	108	112003	111599
1	113	114317	115846
1	114	114522	114316
1	119	118439	118927
1	121	119802	119599
1	125	125688	123967
1	129	128594	129235
1	135	136116	135259
1	136	136558	136298
1	139	139149	139559
1	141	140573	140121
1	143	141738	141412
1	145	142218	142060
1	146	142686	142342
1	154	150528	149074
1	158	153832	153981
1	163	158277	158474

Borrelia burgdorferi - Putative coding regions of novel proteins not similar to know proteins

1	171	168052	166205
1	176	171592	171038
1	186	179607	180089
1	189	182345	182046
1	191	182567	182773
1	199	192561	192716
1	205	196592	197476
1	218	207717	206752
1	219	207733	208437
1	221	209337	208915
1	222	209712	209335
1	231	217179	216025
1	238	223660	223418
1	240	224720	225724
1	242	227006	227275
1	248	231761	231501
1	251	232973	233308
1	252	233669	234004
1	254	235115	235456
1	258	241824	242198
1	261	248009	247773
1	269	256846	255872
1	276	265430	265158
1	279	266582	266298
1	281	268474	268280
1	286	274157	274384
1	292	280495	280274
1	294	281344	281042
1	298	287276	285714
1	303	292943	292644
1	304	293273	293037
1	305	294965	294648
1	308	299427	298699
1	309	299051	299212
1	326	320375	319785
1	327	320425	321036
1	331	324198	324413
1	339	332785	332459
1	341	333503	334138
1	342	334116	334739
1	343	334880	335446
1	350	342916	342443
1	351	344789	342897
1	363	357596	356931
1	367	361065	360859
1	370	362519	362196

Borrelia burgdorferi - Putative coding regions of novel proteins not similar to know proteins

1	374	366905	366114
1	377	368632	369537
1	378	369928	370560
1	379	370532	371353
1	382	375028	373193
1	383	375102	375542
1	387	378677	378198
1	400	394952	394722
1	401	396247	394937
1	403	397569	398327
1	406	399103	399294
1	436	416160	416570
1	445	424660	423950
1	446	425181	424642
1	450	428559	428200
1	451	428933	428619
1	455	432590	431628
1	461	437823	438092
1	463	438690	438313
1	466	440749	440222
1	470	441568	441350
1	471	442039	441614
1	472	442216	442037
1	473	442666	442262
1	476	445202	445017
1	493	462106	462519
1	494	462893	462549
1	504	482111	481035
1	505	481552	481800
1	509	483249	483668
1	512	484864	485157
1	516	489171	488527
1	519	492989	492375
1	520	493626	492997
1	521	494169	494864
1	524	497185	497385
1	525	497674	499254
1	527	500251	501294
1	528	501281	502156
1	558	533912	533667
1	568	541267	541491
1	571	544436	544257
1	572	544565	545068
1	578	549603	551198
1	580	551508	551657
1	581	552337	551513

Borrelia burgdorferi - Putative coding regions of novel proteins not similar to know proteins

1	585	556051	557271
1	590	561342	561139
1	591	561825	561520
1	592	562536	563360
1	596	565758	566519
1	599	568389	568682
1	602	568680	568856
1	605	570829	571167
1	609	576170	577093
1	612	581549	581091
1	614	582910	584013
1	619	589384	588674
1	624	592665	593465
1	626	594542	595405
1	672	631642	632175
1	677	636650	636892
1	678	637059	638078
1	681	640861	640412
1	686	644887	645207
1	689	649716	649961
1	690	650436	650735
1	691	650733	651056
1	693	653303	653689
1	705	664733	664918
1	707	665979	666770
1	718	679155	678391
1	721	680664	681047
1	722	681523	681849
1	724	681809	682171
1	727	682853	683272
1	734	687648	688067
1	739	691613	692290
1	751	707290	707718
1	763	719197	718904
1	764	720030	719257
1	769	722198	722482
1	783	733736	734647
1	785	735554	736618
1	787	737124	739184
1	792	742924	744801
1	799	753128	752655
1	811	766129	765980
1	812	766438	767772
1	815	770062	769790
1	818	771890	772282
1	831	788219	788836

Borrelia burgdorferi - Putative coding regions of novel proteins not similar to know proteins

1	832	788824	789615
1	838	793566	793414
1	840	794295	794119
1	844	796774	796586
1	852	803096	802908
1	858	809371	809970
1	864	816108	816497
1	865	816672	817283
1	866	817281	817838
1	872	823841	824836
1	876	828191	828739
1	877	828749	829147
1	879	831328	831714
1	880	831698	833005
1	885	836201	835677
1	890	841171	840590
1	891	840594	840860
1	899	849453	850148
1	902	851608	852687
1	918	862867	863109
1	920	864292	864705
1	923	865660	865346
1	925	868212	869273
1	928	871012	872580
1	933	878576	879166
1	939	884338	883268
1	940	884999	884325
1	949	892388	892924
1	957	900141	899296
1	958	900534	900139
1	959	901526	900510
1	962	902383	903258

TABLE 4. Borrelia burgdorferi - Putative coding regions of novel proteins similar to know proteins

Contig ID	ORF ID	Start (nt)	Stop (nt)	match acession	match gene name	% sim	% ident
69	1	291	4	gil146582	beta-lactamase [Escherichia coli]	100	98
69	2	692	240	gil344797	galactosidase fusion protein [unidentified]	100	99
26	3	1575	2093	gil458219	ORF 4 [Borrelia burgdorferi]	94	76
2	48	41836	41459	gil47453	ribosomal protein S12 [Streptococcus pneumoniae]	92	85
6	20	14234	12951	bbsl161785	60 kda antigen [Borrelia coriaceae, C053, ATCC 4338, Peptide, 514 aa] [Borrelia coriaceae]	88	67
52	5	1080	1652	gnllPIDle201250	ORF-D gene product [Borrelia burgdorferi]	88	74
152	1	337	26	gnllPIDle158979	orfA gene product [Borrelia burgdorferi]	86	75
71	2	1421	1128	gnllPIDle160437	orfD gene product [Borrelia burgdorferi]	85	46
131	1	381	674	gil458220	ORF 5 [Borrelia burgdorferi]	85	76
3	113	98152	97367	gil1591672	phosphate transport system ATP-binding protein [Methanococcus jannaschii]	84	65
2	107	108403	109485	gil882454	fructose 1,6-bisphosphate aldolase [Escherichia coli]	81	61
19	4	4059	4754	pirlA34520lA34520	29K calcium-binding protein, brain-specific - guinea pig (fragments)	81	56
20	9	6084	5791	gnllPIDle201249	ORF-C gene product [Borrelia burgdorferi]	81	72
2	52	49986	49600	pirlA02771lR7MCML	ribosomal protein L7/L12 - Micrococcus luteus	80	67
14	1	3071	3	gil1522636	M. jannaschii predicted coding region MJECS02 [Methanococcus jannaschii]	80	60
29	2	218	409	gil1752736	gene required for phosphoylation of oligosaccharides/ has high homology with YJR061w [Saccharomyces cerevisiae]	80	37
32	2	719	925	gil433720	CDC25 [Homo sapiens]	80	73
100	1	2	946	gil1522636	M. jannaschii predicted coding region MJECS02 [Methanococcus jannaschii]	80	60

Borrelia burgdorferi - Putative coding regions of novel proteins similar to know proteins

2	106	107148	108239	gnl|PID|e288124	glucose epimerase [Bacillus thuringiensis]	79	66
8	4	4878	4735	gil1543076	outer membrane porin protein Oms28 precursor [Borrelia burgdorferi]	79	69
2	55	51661	51218	gil587583	ribosomal protein L11 [Thermus aquaticus thermophilus]	78	58
4	54	39290	38742	gnl|PID|e160437	orfD gene product [Borrelia burgdorferi]	78	57
5	46	27416	27177	gnl|PID|e253211	ORF YDL065c [Saccharomyces cerevisiae]	78	57
7	4	2382	2966	gnl|PID|e201248	ORF-B gene product [Borrelia burgdorferi]	78	60
19	5	5107	4943	gil882579	CG Site No. 29739 [Escherichia coli]	78	42
78	1	1	171	gnl|PID|e201249	ORF-C gene product [Borrelia burgdorferi]	78	60
105	2	503	742	gnl|PID|e253211	ORF YDL065c [Saccharomyces cerevisiae]	78	57
2	30	24917	23697	gil143795	transfer RNA-Tyr synthetase [Bacillus subtilis]	77	52
6	34	22722	24080	gil466474	cellobiose phosphotransferase enzyme II" [Bacillus stearothermophilus]	77	50
8	1	688	536	gil1017809	similar to dihydropryridine-sensitive l-type, skeletal muscle calcium channel alpha-1 subunit (SP:CIC1_RABIT, P07293) [Caenorhabditis elegans]	77	55
3	91	81071	82183	gil467376	unknown [Bacillus subtilis]	76	58
11	1	208	2	gil1065989	(pos:5995..5997,aa:Met) [Bacillus subtilis]	76	56
68	1	605	3	gnl|PID|e158980	orfC gene product [Borrelia burgdorferi]	76	56
2	9	8488	6674	pir|C30010|C30010	hypothetical ORF-6 protein - Sauroleishmania tarentolae mitochondrion (SGC6)	75	50
2	37	31639	32163	gil1573470	H. influenzae predicted coding region HI0491 [Haemophilus influenzae]	75	55
2	56	52261	51701	gil396321	nusG [Escherichia coli]	75	56

Borrelia burgdorferi - Putative coding regions of novel proteins similar to know proteins

20	1	653	414	gil520778	protein p23 [Borrelia burgdorferi]	75	62
20	3	2437	1652	gnl|PID|e201249	ORF-C gene product [Borrelia burgdorferi]	75	60
58	1	856	62	gnl|PID|e201249	ORF-C gene product [Borrelia burgdorferi]	75	37
68	3	1153	578	gil458217	ORF 2 [Borrelia burgdorferi]	75	55
117	1	744	388	gil520783	unknown [Borrelia burgdorferi]	75	42
130	1	1	684	gnl|PID|e160436	orfA gene product [Borrelia burgdorferi]	75	58
2	36	30506	31693	gil1020317	S-adenosylmethionine synthetase [Staphylococcus aureus]	74	57
2	109	111301	109871	gil396501	aspartyl-tRNA synthetase [Thermus aquaticus thermophilus]	74	52
3	101	92143	91103	gil1651962	hypothetical protein [Synechocystis sp.]	74	49
20	5	4080	2974	gnl|PID|e158979	orfA gene product [Borrelia burgdorferi]	74	56
36	2	468	1253	gnl|PID|e158984	orfC gene product [Borrelia burgdorferi]	74	59
42	1	396	719	gil1655798	CdsK [Borrelia burgdorferi]	74	58
2	10	6810	7022	gil406135	glycoprotein 120 [Simian immunodeficiency virus]	73	53
2	29	23695	21395	gil511145	hemolysin [Serpulina hyodysenteriae]	73	52
3	56	44789	44262	gil1595810	type-I signal peptidase SpsB [Staphylococcus aureus]	73	47
3	73	64881	62341	gnl|PID|e268456	unknown [Mycobacterium tuberculosis]	73	54
3	100	89800	91113	gil500705	Similar to Seryl-tRNA synthetase [Saccharomyces cerevisiae]	73	56
3	106	92803	93513	gnl|PID|e243681	ORF YGR248w [Saccharomyces cerevisiae]	73	63
4	4	3697	3512	gil562035	NADH dehydrogenase, subunit 5 [Acanthamoeba castellanii]	73	53
7	9	8519	8079	gil694092	emml gene product [Streptococcus pyogenes]	73	40

Borrelia burgdorferi - Putative coding regions of novel proteins similar to know proteins

8	16	17562	17756	gil1500401	reverse gyrase [Methanococcus jannaschii]	73	40
14	3	4280	4438	gil520778	protein p23 [Borrelia burgdorferi]	73	55
19	9	7074	6742	gil1773311	NADH dehydrogenase [Ceanothus cuneatus]	73	36
25	3	2369	2587	gil1655790	CdsC [Borrelia burgdorferi]	73	64
78	2	176	619	gnllPIDle2012 50	ORF-D gene product [Borrelia burgdorferi]	73	50
108	1	2	382	gil1573074	adhesin B precursor (fimA) [Haemophilus influenzae]	73	41
120	1	97	342	gil1978	heat shock protein 70 [Sus scrofa]	73	46
3	64	51644	54013	gil1574437	sporulation protein (spoIIIE) [Haemophilus influenzae]	72	51
5	6	2899	2654	gil212383	myosin heavy chain [Gallus gallus]	72	41
6	31	22140	21799	gil895748	putative cellobiose phosphotransferase enzyme II' [Bacillus subtilis]	72	46
8	8	8812	9600	gil1655859	Orf1 [Borrelia hermsii]	72	55
10	12	8579	8376	gil536681	ORF YBR257w [Saccharomyces cerevisiae]	72	36
45	2	1440	394	gil1699017	ErpB2 [Borrelia burgdorferi]	72	42
2	2	1342	2796	gil285623	pyruvate kinase [Bacillus stearothermophilus]	71	52
2	31	26272	24911	pirlS58522lS5 8522	glycyl-tRNA synthetase - Thermus thermophilus	71	54
2	64	60156	58684	gil459009	similar to multifunctional aminoacyl-tRNA synthetase, especially to the prolyl-tRNA synthetase region [Caenorhabditis elegans]	71	48
3	66	55240	54275	gil217121	ORF1 [Synechococcus elongatus]	71	52
3	104	92345	92175	gil44228	secretion protein SecY (AA 1-482) [Mycoplasma capricolum]	71	42
5	43	25567	25734	gil213778	sodium-hydrogen exchange protein-beta [Oncorhynchus mykiss]	71	50
7	3	1179	2384	gil458216	ORF 1 [Borrelia burgdorferi]	71	60
20	4	2964	2392	gil458217	ORF 2 [Borrelia burgdorferi]	71	47
51	2	984	2066	gil1373144	ErpD [Borrelia burgdorferi]	71	41
54	1	251	883	gil145280	ORF1 [Escherichia coli]	71	40

Borrelia burgdorferi - Putative coding regions of novel proteins similar to know proteins

2	54	51233	50538	splQ06797lRL1_BACSU	50S RIBOSOMAL PROTEIN L1 (BL1).	70	48
2	116	114025	113744	gil1673757	(AE000012) Mycoplasma pneumoniae, phosphocarrier protein HPr; similar to GenBank Accession Number A49683, from M. capricolum [Mycoplasma pneumoniae]	70	41
3	4	1684	2220	gil153906	CheW protein [Salmonella typhimurium]	70	48
3	84	74775	73225	gnllPIDle283919	glycerol kinase [Sulfolobus solfataricus]	70	60
3	107	93500	93273	gil836815	cdc4 gene product which is essential for initiation of DNA replication in yeast [Saccharomyces cerevisiae]	70	35
4	1	926	123	gil167913	Thy1 protein [Dictyostelium discoideum]	70	51
4	47	35616	35807	gil48808	dciAE gene product [Bacillus subtilis]	70	58
4	65	48320	47976	gil1421734	ORF 5 [Borrelia burgdorferi]	70	48
6	23	16458	15904	gil1655860	Orf2 [Borrelia hermsii]	70	54
17	4	2940	3173	gil1255880	F01G12.6 gene product [Caenorhabditis elegans]	70	40
20	8	5470	5237	gil1236921	Var1p [Saccharomyces douglasii]	70	47
23	5	4173	3970	pirlS16447lS16447	NADH dehydrogenase (ubiquinone) (EC 1.6.5.3) chain 4 - wheat mitochondrion	70	47
36	3	1270	1653	gnllPIDle160437	orfD gene product [Borrelia burgdorferi]	70	50
2	69	65752	63860	gil151932	fructose enzyme II [Rhodobacter capsulatus]	69	42
3	114	99712	98150	gil1303856	YqgI [Bacillus subtilis]	69	46
4	36	25614	24694	gil1663561	orfI; product unknown [Borrelia burgdorferi]	69	46
6	21	14584	14204	gil1616644	P30 [Borrelia burgdorferi]	69	47
12	12	7025	7258	gil150176	protein 69 [Mycoplasma hyorhinis]	69	38
12	14	8414	8587	gil13233	ND6 (AA 1 - 296) [Podospora anserina]	69	26
54	2	1332	2402	gnllPIDle158979	orfA gene product [Borrelia burgdorferi]	69	46
2	35	29769	30518	gil473817	'ORF' [Escherichia coli]	68	42
2	79	72330	72980	gil1498049	adenylate kinase [Paracoccus denitrificans]	68	37

Borrelia burgdorferi - Putative coding regions of novel proteins similar to know proteins

2	104	104748	106385	gil1574032	hypothetical [Haemophilus influenzae]	68	42
3	78	68895	68287	gnl|PID|e255117	hypothetical protein [Bacillus subtilis]	68	51
3	98	88992	86074	gil927711	D9461.18p; CAI: 0.15 [Saccharomyces cerevisiae]	68	52
3	111	96519	97364	gil1707057	coded for by C. elegans cDNA CEESS55F; coded for by C. elegans cDNA yk84a1.3; coded for by C. elegans cDNA yk78g7.3; coded for by C. elegans cDNA yk168g9.5; coded for by C. elegans cDNA yk78g7.5; coded for by C. elegans cDNA yk84a1.5; strong s	68	52
4	56	40648	40046	gil458217	ORF 2 [Borrelia burgdorferi]	68	54
4	57	41916	40678	gil458216	ORF 1 [Borrelia burgdorferi]	68	57
6	24	17296	16520	gil1655859	Orf1 [Borrelia hermsii]	68	43
7	5	2894	3694	gil1655859	Orf1 [Borrelia hermsii]	68	48
29	6	3832	3254	gil458217	ORF 2 [Borrelia burgdorferi]	68	46
72	2	927	1133	gil577175	L8479.4 gene product [Saccharomyces cerevisiae]	68	44
2	57	52752	52558	gil1001264	50S ribosomal protein L33 [Synechocystis sp.]	67	56
3	65	54290	54051	gil710340	ribosomal protein S21 [Myxococcus xanthus]	67	49
3	79	69068	70114	gil460955	TagE [Vibrio cholerae]	67	38
3	81	70653	71150	gil467420	unknown [Bacillus subtilis]	67	42
3	110	94703	96502	gnl|PID|e267607	alanyl-tRNA synthatase [Thermus aquaticus thermophilus]	67	51
4	42	30304	31941	bbs|161785	60 kda antigen [Borrelia coriaceae, C053, ATCC 4338, Peptide, 514 aa] [Borrelia coriaceae]	67	49
12	6	3590	2967	gnl|PID|e160437	orfD gene product [Borrelia burgdorferi]	67	41
12	9	5524	6276	gil1655859	Orf1 [Borrelia hermsii]	67	51
12	10	6611	6889	gnl|PID|e8903	SERA protein [Plasmodium falciparum]	67	48
17	6	4995	5906	gil1752736	gene required for phosphoylation of oligosaccharides/ has high homology with YJR061w [Saccharomyces cerevisiae]	67	37
34	2	1221	1817	gnl|PID|e1589	orfB gene product [Borrelia burgdorferi]	67	47

Borrelia burgdorferi - Putative coding regions of novel proteins similar to know proteins

				83			
58	2	1347	796	gil458217	ORF 2 [Borrelia burgdorferi]	67	52
2	33	28572	27751	gil340613	A 'c' was inserted after nt 369 (=nt 10459 in genomic sequence (M10126)) to correct -1 frameshift probably due to gel compression [Leishmania tarentolae]	66	40
2	73	69021	69908	gil153903	methyltransferase (cheR; EC 2.1.1.24) [Salmonella typhimurium]	66	42
2	93	93739	94524	gil45713	P.putida genes rpmH, rnpA, 9k, 60k, 50k, gidA, gidB, uncI and uncB [Pseudomonas putida]	66	41
3	9	6009	6902	gnl|PID|e263931	OrfD [Streptococcus pneumoniae]	66	47
4	28	20922	20665	gil471731	vacuolating cytotoxin homolog [Helicobacter pylori]	66	50
4	64	47985	47107	gil1421735	ORF 6 [Borrelia burgdorferi]	66	43
6	13	7227	8591	gil1591045	hypothetical protein (SP:P31466) [Methanococcus jannaschii]	66	48
34	4	2556	3161	gil458218	ORF 3 [Borrelia burgdorferi]	66	42
37	1	982	689	gil974334	non-receptor tyrosine kinase [Dictyostelium discoideum]	66	55
3	77	68191	66395	gil1651216	Pz-peptidase [Bacillus licheniformis]	65	47
3	123	105911	104070	gil1575784	DNA mismatch repair protein [Aquifex pyrophilus]	65	45
6	9	5726	7126	gil1591045	hypothetical protein (SP:P31466) [Methanococcus jannaschii]	65	49
8	9	9684	10325	gnl|PID|e201250	ORF-D gene product [Borrelia burgdorferi]	65	48
10	1	3	971	gil1373144	ErpD [Borrelia burgdorferi]	65	47
13	5	3956	3411	gil1209872	REV [Borrelia burgdorferi]	65	47
2	76	70509	71069	pir|A00547|XYEBET	protein-glutamate methylesterase (EC 3.1.1.61) - Salmonella typhimurium	64	45
3	61	48610	50838	gil1001335	soluble lytic transglycosylase [Synechocystis sp.]	64	42
4	5	3519	3773	gil1263021	M protein [Streptococcus pyogenes]	64	32
4	53	38288	37824	gil1373141	ORF-10 [Borrelia burgdorferi]	64	50

Borrelia burgdorferi - Putative coding regions of novel proteins similar to know proteins

6	10	5985	5824	gil40271	delta-endotoxin CryIG protoxin [Bacillus thuringiensis]	64	30
7	7	7798	4499	gil1041785	rhoptry protein [Plasmodium yoelii]	64	35
7	30	19738	19289	gil1209840	2.9-3 ORF-D [Borrelia burgdorferi]	64	46
11	3	1608	2339	gil1652934	hypothetical protein [Synechocystis sp.]	64	30
16	1	537	839	gnl|PID|e276380	AARP1 protein [Plasmodium falciparum]	64	44
19	1	308	1177	gil1553115	P35 antigen protein [Borrelia burgdorferi]	64	35
42	3	1928	1788	gil1752736	gene required for phosphoylation of oligosaccharides/ has high homology with YJR061w [Saccharomyces cerevisiae]	64	41
142	1	589	2	gil162142	kinetoplast-associated protein [Trypanosoma cruzi]	64	52
2	3	2837	2592	gnl|PID|e236274	ZK287.2 [Caenorhabditis elegans]	63	27
2	15	12750	11320	gil1652577	carboxyl-terminal protease [Synechocystis sp.]	63	49
2	32	27753	26266	sp|P15189|SYE_RHIME	GLUTAMYL-TRNA SYNTHETASE (EC 6.1.1.17) (GLUTAMATE--TRNA LIGASE) (GLURS).	63	48
2	77	71067	72308	gil1041116	TRAB [Plasmid pPD1]	63	34
3	2	1056	58	gil1098641	Bts1p [Saccharomyces cerevisiae]	63	43
3	82	71398	71237	gil1339938	EC 1.1.99.5 [Mus musculus]	63	47
3	83	72845	71349	gil763191	glycerol 3 phosphate dehydrogenase [Saccharomyces cerevisiae]	63	37
3	85	75552	74773	gil142997	glycerol uptake facilitator [Bacillus subtilis]	63	45
7	6	3747	4304	gnl|PID|e201250	ORF-D gene product [Borrelia burgdorferi]	63	40
7	38	24123	24956	gil467330	replicative DNA helicase [Bacillus subtilis]	63	40
11	5	4161	3853	gil1592217	bifunctional protein [Methanococcus jannaschii]	63	38
12	13	9558	7906	gil633167	adenine deaminase [Bacillus subtilis]	63	48
32	1	753	268	gil520783	unknown [Borrelia burgdorferi]	63	42
2	68	63866	62745	gil146722	phosphomannose isomerase [Escherichia coli]	62	45
2	75	69920	70573	gil145524	cheB peptide [Escherichia coli]	62	28

2	95	96334	95492	gil40031	spoOJ93 gene product [Bacillus subtilis]	62	36
3	67	57341	55212	gil1574144	single-stranded-DNA-specific exonuclease (recJ) [Haemophilus influenzae]	62	40
3	76	66414	65677	gil1477770	unknown [Helicobacter pylori]	62	37
6	1	1762	104	gil1072419	glcB gene product [Staphylococcus carnosus]	62	43
18	4	4431	5144	gil1591493	glutamine transport ATP-binding protein Q [Methanococcus jannaschii]	62	36
19	8	6743	6976	gil1513302	CigB [Dictyostelium discoideum]	62	56
20	6	4563	4378	bbsl144872	Fu=putative serine/threonine kinase [Drosophila melanogaster, Peptide Partial Mutant, 152 aa] [Drosophila melanogaster]	62	37
81	1	56	538	gnllPIDle153957	ORF-A gene product [Borrelia burgdorferi]	62	36
106	2	586	356	gil1151158	repeat organellar protein [Plasmodium chabaudi]	62	43
114	1	138	629	gnllPIDle153957	ORF-A gene product [Borrelia burgdorferi]	62	36
2	117	114352	114032	gil173128	ubiquitin-specific processing protease [Saccharomyces cerevisiae]	61	32
3	55	42737	44236	gil143999	dnaK homologue [Borrelia burgdorferi]	61	41
3	57	44821	46083	gil1653709	lipoprotein NlpD [Synechocystis sp.]	61	50
3	125	110052	109261	gil1303863	YqgP [Bacillus subtilis]	61	45
4	63	47119	46478	gil1421736	ORF 7 [Borrelia burgdorferi]	61	34
7	35	21496	22971	gil1655797	CdsJ [Borrelia burgdorferi]	61	44
8	7	8300	8872	gil458217	ORF 2 [Borrelia burgdorferi]	61	48
12	8	5006	5551	gil458217	ORF 2 [Borrelia burgdorferi]	61	50
14	10	9398	8652	gnllPIDle201250	ORF-D gene product [Borrelia burgdorferi]	61	44
15	12	6206	4377	gil836624	methyltransferase [Bacillus aneurinolyticus]	61	38
16	4	2449	2240	gil1066497	Similar to S. cerevisiae hypothetical protein Ykl012p (Swiss Prot. accession number P33203) and C. elegans hypothetical protein ZK1098.1 (Swiss Prot. accession number P34600) [Saccharomyces	61	38

Borrelia burgdorferi - Putative coding regions of novel proteins similar to know proteins

					cerevisiae]		
29	4	2323	1853	gnl|PID|e160437	orfD gene product [Borrelia burgdorferi]	61	45
50	2	1374	1156	gnl|PID|e276380	AARP1 protein [Plasmodium falciparum]	61	52
2	18	14371	15369	gil1573923	prolipoprotein diacylglyceryl transferase (lgt) [Haemophilus influenzae]	60	57
3	118	101571	100690	gil1001260	hypothetical protein [Synechocystis sp.]	60	47
3	120	101692	102273	gil1399829	elongation factor P [Synechococcus PCC7942]	60	34
6	32	21869	22162	gil192960	L-type calcium channel alpha-1 [Mus musculus]	60	50
7	37	23373	24101	gil458217	ORF 2 [Borrelia burgdorferi]	60	40
8	11	13570	13851	gil1065989	(pos:5995..5997,aa:Met) [Bacillus subtilis]	60	47
14	5	5327	5091	gil147158	pfs [Escherichia coli]	60	51
15	7	3316	2984	gil153727	M protein [group G streptococcus]	60	36
27	3	2744	3772	pirlS40422lS40422	hypothetical protein - Staphylococcus aureus	60	31
2	62	57446	58672	gil143002	proton glutamate symport protein [Bacillus caldotenax]	59	34
2	82	74989	74051	gil1651878	regulatory components of sensory transduction system [Synechocystis sp.]	59	38
2	89	92119	91322	gil467425	unknown [Bacillus subtilis]	59	38
2	92	93010	93663	pirlA30191lA30191	hypothetical protein L - Bacillus subtilis (fragment)	59	39
2	118	115604	114315	gil39269	sigma factor (ntrA) (AA 1-502) [Azotobacter vinelandii]	59	35
4	41	29875	29210	gil1209831	lipoprotein [Borrelia burgdorferi]	59	34
6	4	3323	2058	gil624056	contains 4 ankyrin repeats; similar to D. melanogaster notch protein, Swiss-Prot Accession Number P07027 [Paramecium bursaria Chlorella virus 1]	59	37
6	25	17793	17257	gnl|PID|e201248	ORF-B gene product [Borrelia burgdorferi]	59	43

6	33	22493	22125	gil153677	enzyme III [Streptococcus mutans]	59	36
10	9	6241	6026	gil1339977	skeletal myosin heavy chain [Thunnus thynnus]	59	40
19	6	5383	5970	gil160299	glutamic acid-rich protein [Plasmodium falciparum]	59	30
25	7	4008	3742	gil1055144	similar to galactoside 3(4)-l-fucosyltransferase [Caenorhabditis elegans]	59	37
59	1	835	8	gil1359436	Mag44 [Dermatophagoides farinae]	59	24
2	27	21414	20317	gil974332	NAD(P)H-dependent dihydroxyacetone-phosphate reductase [Bacillus subtilis]	58	41
3	5	2156	3049	gil1653618	hypothetical protein [Synechocystis sp.]	58	35
3	109	93822	94718	gil790935	fliG [Treponema denticola]	58	31
4	3	2423	3340	gil1553115	P35 antigen protein [Borrelia burgdorferi]	58	39
4	35	24696	24238	gil1663562	orfII; product unknown [Borrelia burgdorferi]	58	32
4	46	35509	34904	gnllPIDle264708	myosin heavy chain [Sus scrofa]	58	31
15	8	3683	3468	gil457336	Pv200 [Plasmodium vivax]	58	32
50	3	1941	1498	gnllPIDle220350	brca2 gene product [Homo sapiens]	58	41
55	1	2322	247	gil1522636	M. jannaschii predicted coding region MJECS02 [Methanococcus jannaschii]	58	32
2	53	50563	50045	gil786163	Ribosomal Protein L10 [Bacillus subtilis]	57	29
3	117	100606	99710	gil1303855	YqgH [Bacillus subtilis]	57	30
6	38	26564	26232	gil1499632	M. jannaschii predicted coding region MJ0809 [Methanococcus jannaschii]	57	40
8	10	12350	13117	gil1553115	P35 antigen protein [Borrelia burgdorferi]	57	33
11	4	3183	2470	pirlA45605lA45605	mature-parasite-infected erythrocyte surface antigen MESA - Plasmodium falciparum	57	35
14	8	7117	7899	gil1553115	P35 antigen protein [Borrelia burgdorferi]	57	36
15	6	3027	2818	gnllPIDle261409	nuclear/mitotic apparatus protein [Xenopus laevis]	57	30
17	1	336	1178	gil473817	'ORF' [Escherichia coli]	57	31
20	2	1654	1064	gnllPIDle201250	ORF-D gene product [Borrelia burgdorferi]	57	34

Borrelia burgdorferi - Putative coding regions of novel proteins similar to know proteins

2	111	113765	111636	gil148316	NaH-antiporter protein [Enterococcus hirae]	56	32
3	80	70112	70669	gil1372995	OrfH [Borrelia burgdorferi]	56	24
3	116	98976	99212	pirlE22845lE22845	hypothetical protein 4 - Trypanosoma brucei mitochondrion (SGC6)	56	36
6	26	18732	17791	gil1655797	CdsJ [Borrelia burgdorferi]	56	41
7	21	14706	13510	gil1574247	H. influenzae predicted coding region HI1410 [Haemophilus influenzae]	56	32
11	8	6722	7087	gnllPIDle242897	aBIM [Lactococcus lactis]	56	28
53	7	2446	2018	gil1421737	ORF 8 [Borrelia burgdorferi]	56	38
61	2	712	1410	gil583161	albumin binding protein [unidentified]	56	35
2	6	3866	3573	gil290487	50S ribosomal subunit protein L28 [Escherichia coli]	55	37
2	14	11322	10585	gil1303811	YqeU [Bacillus subtilis]	55	33
2	34	28640	29782	gil558266	orf gene product [Wolinella succinogenes]	55	30
2	71	66669	67415	gil397486	endonuclease G [Bos taurus]	55	33
3	87	75924	76550	gil403984	deoxyguanosine kinase/deoxyadenosine kinase(I) subunit [Lactobacillus acidophilus]	55	38
4	66	48434	48958	gil1100900	70 kDa heat shock protein [Theileria parva]	55	32
140	1	322	68	gil15611	gene 17, tail fiber protein [Bacteriophage T7]	55	38
4	34	24244	23867	gil1663563	orfIII; product unknown [Borrelia burgdorferi]	54	31
5	9	5510	4179	gil1513238	ORFveg132; similar to Caenorhabditis elegans ORF F59B10.1 encoded by EMBL Accession Number Z49132 [Dictyostelium discoideum]	54	25
5	45	27187	25895	gnllPIDle261409	nuclear/mitotic apparatus protein [Xenopus laevis]	54	30
7	28	17905	18162	gil36501	C protein [Homo sapiens]	54	41
11	6	4415	5215	gil1707287	putative outer membrane protein [Borrelia burgdorferi]	54	25
19	2	1674	2501	gil392799	G5/D6 ORF [Dictyostelium discoideum]	54	25
29	5	3284	2532	gnllPIDle158980	orfC gene product [Borrelia burgdorferi]	54	33
31	3	3328	4137	pirlS41649lS4	DNA polymerase - Plasmodium falciparum	54	28

Borrelia burgdorferi - Putative coding regions of novel proteins similar to know proteins

				1649			
32	5	2560	2865	gil499695	bud-emergence protein [Saccharomyces cerevisiae]	54	36
95	1	95	997	gil763227	Rpi1p [Saccharomyces cerevisiae]	54	37
2	16	13235	14383	gil1165254	YlxH [Borrelia burgdorferi]	53	33
2	72	68814	68179	gil940842	orf 06111 gene product [Saccharomyces cerevisiae]	53	28
3	3	1032	1646	gil1592021	cell division protein J [Methanococcus jannaschii]	53	32
4	18	14627	14427	gil1039462	vacuolar aspartic proteinase precursor [Candida albicans]	53	35
5	63	34850	34152	gil1699017	ErpB2 [Borrelia burgdorferi]	53	25
10	5	3672	3893	gil398581	XLR related protein [Mus musculus]	53	27
15	17	8485	8925	gnl|PID|e248324	T07C12.4 [Caenorhabditis elegans]	53	28
25	5	3497	3679	gil1055100	coded for by C. elegans cDNA yk54h9.5; coded for by C. elegans cDNA yk54h9.3; similar to matrin F/G (DNA binding protein, SP:MAFG_RAT, Q00910) [Caenorhabditis elegans]	53	25
29	1	70	291	gil398581	XLR related protein [Mus musculus]	53	27
34	3	1787	2527	gil1655859	Orf1 [Borrelia hermsii]	53	42
2	1	3	1265	gil496254	fibronectin/fibrinogen-binding protein [Streptococcus pyogenes]	52	29
2	110	111638	111276	gil1573287	aspartyl-tRNA synthetase (aspS) [Haemophilus influenzae]	52	35
4	8	5323	6150	gil457146	rhoptry protein [Plasmodium yoelii]	52	24
4	44	32562	31999	gil1151158	repeat organellar protein [Plasmodium chabaudi]	52	26
7	29	18485	18808	gnl|PID|e243927	ORF YGR023w [Saccharomyces cerevisiae]	52	26
25	4	3287	3499	gil500655	YHR146w gene product [Saccharomyces cerevisiae]	52	35
92	2	38	241	gil1236411	NADH dehydrogenase, subunit 5 [Allomyces macrogynus]	52	52
148	2	119	322	gil1236411	NADH dehydrogenase, subunit 5 [Allomyces macrogynus]	52	52
2	120	116131	115577	gil1591434	chromate resistance protein A [Methanococcus	51	29

Borrelia burgdorferi - Putative coding regions of novel proteins similar to know proteins

					jannaschii]		
4	9	6362	7153	gil1553115	P35 antigen protein [Borrelia burgdorferi]	51	26
10	10	6603	7196	gnl|PID|e256393	anti-P.falciparum antigenic polypeptide [Saimiri sciureus]	51	34
11	12	10333	9422	pir|A42771|A42771	reticulocyte-binding protein 1 - Plasmodium vivax	51	31
19	7	5919	6179	gil173241	ZIP1 protein [Saccharomyces cerevisiae]	51	38
23	1	3	287	gil1498320	cell wall-associated protease precursor [Bacillus subtilis]	51	25
2	105	106383	107126	gil580905	B.subtilis genes rpmH, rnpA, 50kd, gidA and gidB [Bacillus subtilis]	50	32
3	1	1	195	gnl|PID|e220201	rps5 gene product [Plasmodium falciparum]	50	38
3	62	50808	51653	gil882579	CG Site No. 29739 [Escherichia coli]	50	31
3	119	100766	101014	gil1086864	T03G11.2 gene product [Caenorhabditis elegans]	50	39
4	32	23555	22992	gil1663565	orfV; product unknown [Borrelia burgdorferi]	50	36
5	8	4168	3470	gil49402	M1.1 protein [Streptococcus pyogenes]	50	27
10	7	5190	4612	gnl|PID|e158981	orfE gene product [Borrelia burgdorferi]	50	28
11	2	1277	504	gil1553115	P35 antigen protein [Borrelia burgdorferi]	50	26
13	3	1948	1634	gnl|PID|e268243	p21 [Borrelia afzelii]	50	32
92	3	582	941	gnl|PID|e201250	ORF-D gene product [Borrelia burgdorferi]	50	40
148	1	339	4	gnl|PID|e236901	unknown [Saccharomyces cerevisiae]	50	34
28	3	2001	2630	gil499325	STARP antigen [Plasmodium falciparum]	49	22
3	10	6881	7180	gil156218	putative [Caenorhabditis elegans]	48	32
3	75	65683	65066	gil1574476	dedA protein (dedA) [Haemophilus influenzae]	48	22
3	112	97006	96743	gil915207	gastric mucin [Sus scrofa]	48	27
7	23	14743	14970	gil172294	protein-tyrosine phosphatase [Saccharomyces cerevisiae]	48	33

Borrelia burgdorferi - Putative coding regions of novel proteins similar to know proteins

11	11	7980	9293	gil1046137	M. genitalium predicted coding region MG422 [Mycoplasma genitalium]	48	28
28	4	2628	2825	gil1591322	chorismate mutase subunit B [Methanococcus jannaschii]	48	30
2	8	5526	6677	gnl|PID|e220245	frameshift [Plasmodium falciparum]	47	27
2	60	55075	55803	gil710551	ankyrin 3 [Mus musculus]	47	29
2	94	94515	95240	gil1592264	type I restriction enzyme [Methanococcus jannaschii]	47	34
4	11	9057	9941	gil1553115	P35 antigen protein [Borrelia burgdorferi]	47	32
2	12	9986	9471	gil1022328	Four tandem repeats of a DNA-binding domain known as the AT-hook are found at the carboxy terminus of CarD. This protein has been purified and found to bind in vitro to a promoter region [Myxococcus xanthus]	46	28
3	89	78904	77324	gil1573271	apolipoprotein N-acyltransferase (cute) [Haemophilus influenzae]	46	32
6	36	24361	25719	gil1592272	ribosomal protein S19 [Methanococcus jannaschii]	46	23
10	13	9895	8816	gil160299	glutamic acid-rich protein [Plasmodium falciparum]	46	21
13	4	3412	3648	gnl|PID|e275506	C41G6.i [Caenorhabditis elegans]	46	37
138	1	632	15	gil157006	bicaudalD protein [Drosophila melanogaster]	46	23
3	124	109271	105909	gil1499043	M. jannaschii predicted coding region MJ0263 [Methanococcus jannaschii]	45	23
4	17	14212	15465	pir|S30782|S30782	integrin homolog - yeast (Saccharomyces cerevisiae)	45	23
23	4	3950	4852	gnl|PID|e236901	unknown [Saccharomyces cerevisiae]	45	27
92	1	258	4	gnl|PID|e236901	unknown [Saccharomyces cerevisiae]	45	27
3	90	79020	81044	gnl|PID|e236483	F54G8.4 [Caenorhabditis elegans]	44	31
12	7	4075	5019	gil1151158	repeat organellar protein [Plasmodium chabaudi]	44	26

17	3	1735	2142	pirlA42771lA42771	reticulocyte-binding protein 1 - Plasmodium vivax	44	26
22	7	4179	2827	gil563812	XCAP-C [Xenopus laevis]	44	20
31	2	1682	2761	gil1438951	cutinase negative acting protein [Fusarium solani f. sp. pisi]	43	23
3	7	4086	5186	gil343962	VAR1 protein [Candida glabrata]	42	25
28	1	110	496	gil157804	laminin B2 chain [Drosophila melanogaster]	42	23
28	5	2889	3833	pirlS30782lS30782	integrin homolog - yeast (Saccharomyces cerevisiae)	42	18
34	1	209	1234	gil1655797	CdsJ [Borrelia burgdorferi]	42	27
65	3	1035	1415	gil1654220	variable major protein 16 [Borrelia hermsii]	42	34
2	11	9544	8486	gnllPIDle16326	MURF2 protein (AA 1-348) [Crithidia fasciculata]	41	26
3	122	104072	103017	gil1151158	repeat organellar protein [Plasmodium chabaudi]	41	20
18	6	5122	6366	gil1591494	M. jannaschii predicted coding region MJ0797 [Methanococcus jannaschii]	40	20
6	6	4662	3964	gil600448	var1 protein (aa 1-339) [Candida utilis]	39	24
4	10	7637	8914	gil1293695	microfilarial sheath protein SHP3 [Litomosoides sigmodontis]	37	19

TABLE 5. Borrelia burgdorferi - Coding regions containing to know proteins

Contig ID	Orf ID	Start (nt)	Stop (nt)	match acession	match gene name	percent ident	HSP nt length
2	20	15372	17402	gb|M90084|	Borrelia burgdorferi 22 kD antigen	100	786
2	21	16672	16310	gb|M90084|	Borrelia burgdorferi 22 kD antigen	100	56
2	22	17362	17099	gb|M90084|	Borrelia burgdorferi 22 kD antigen	100	264
2	23	17415	17876	emb|X70826|BBLA7	B.burgdorferi gene for lipoprotein	100	57
2	24	18522	17923	emb|X70826|BBLA7	B.burgdorferi gene for lipoprotein	100	600
2	25	18606	20009	emb|X78708|BBYSC1	B.bergdorferi (ZS7) YSC1-like gene	100	1404
2	26	19981	20295	emb|X78708|BBYSC1	B.bergdorferi (ZS7) YSC1-like gene	99	314
2	38	32899	32174	gb|U49938|	Borrelia burgdorferi potential virulence gene cluster membrane proteins BmpC (bmpC) and BmpA (bmpA), BmpB protein (bmpB), putative protein 4, Mg ion transporter MgtE (mgtE), protein kinase C1 inhibitor PKCI (pkci) genes, complete cds	98	130
2	39	33315	32863	gb|U49938|	Borrelia burgdorferi potential virulence gene cluster membrane proteins BmpC (bmpC) and BmpA (bmpA), BmpB protein (bmpB), putative protein 4, Mg ion transporter MgtE (mgtE), protein kinase C1 inhibitor PKCI (pkci) genes, complete cds	100	453
2	40	34718	33333	gb|U49938|	Borrelia burgdorferi potential virulence gene cluster membrane proteins BmpC (bmpC) and BmpA (bmpA), BmpB protein (bmpB), putative protein 4, Mg ion transporter MgtE (mgtE), protein kinase C1 inhibitor PKCI (pkci) genes, complete cds	99	1386
2	41	36211	34751	gb|U49938|	Borrelia burgdorferi potential virulence gene cluster membrane proteins BmpC (bmpC) and BmpA (bmpA), BmpB protein (bmpB), putative protein 4, Mg ion transporter MgtE (mgtE), protein kinase C1 inhibitor PKCI (pkci) genes, complete cds	99	1461
2	42	36899	36288	gb|L24194|	Borrelia burgdorferi immunodominant antigen P39	99	606

Borrelia burgdorferi - Coding regions containing to know proteins

					gene, complete cds		
2	43	37335	36811	gb|L35050|	Borrelia burgdorferi (clone pB46) membrane lipoprotein A (bmpA) gene, 3' end, membrane lipoprotein (bmpB) gene, 5' end	98	457
2	44	38426	37401	gb|L24194|	Borrelia burgdorferi immunodominant antigen P39 gene, complete cds	99	1026
2	45	39595	38462	gb|U49938|	Borrelia burgdorferi potential virulence gene cluster membrane proteins BmpC (bmpC) and BmpA (bmpA), BmpB protein (bmpB), putative protein 4, Mg ion transporter MgtE (mgtE), protein kinase C1 inhibitor PKCI (pkci) genes, complete cds	99	1134
2	46	40947	39838	gb|U35450|	Borrelia burgdorferi membrane protein D (bmpD) gene, complete cds	99	1110
2	47	41461	40961	gb|U35450|	Borrelia burgdorferi membrane protein D (bmpD) gene, complete cds	100	82
2	49	46052	41901	gb|L48488|	Borrelia burgdorferi RNA polymerase beta subunit (rpoB) gene, complete cds, RNA polymerase beta' subunit (rpoC) gene, 5' end of cds	97	76
2	51	49535	46050	gb|L48488|	Borrelia burgdorferi RNA polymerase beta subunit (rpoB) gene, complete cds, RNA polymerase beta' subunit (rpoC) gene, 5' end of cds	98	2490
2	83	79470	74977	gb|U03396|	Borrelia burgdorferi B31 Ala-tRNA (alaT), Ile-tRNA (ileT), 16S rRNA, 23S rRNA (rrlA and rrlB), and 5S rRNA (rrfA and rrfB) genes, complete sequence	97	131
2	84	84351	84620	gb|M88330|	Borrelia burgdorferi 23S ribosomal RNA gene	100	270
2	85	86923	86066	gb|U03396|	Borrelia burgdorferi B31 Ala-tRNA (alaT), Ile-tRNA (ileT), 16S rRNA, 23S rRNA (rrlA and rrlB), and 5S rRNA (rrfA and rrfB) genes, complete sequence	95	386
2	86	87637	87041	gb|U03396|	Borrelia burgdorferi B31 Ala-tRNA (alaT), Ile-tRNA (ileT), 16S rRNA, 23S rRNA (rrlA and rrlB), and 5S rRNA (rrfA and rrfB) genes, complete sequence	99	209

2	87	88424	88116	gb|U03396|	Borrelia burgdorferi B31 Ala-tRNA (alaT), Ile-tRNA (ileT), 16S rRNA, 23S rRNA (rrlA and rrlB), and 5S rRNA (rrfA and rrfB) genes, complete sequence	96	210
2	88	91249	90680	gb|U03396|	Borrelia burgdorferi B31 Ala-tRNA (alaT), Ile-tRNA (ileT), 16S rRNA, 23S rRNA (rrlA and rrlB), and 5S rRNA (rrfA and rrfB) genes, complete sequence	100	570
2	96	98846	96393	emb|Z12165|BBGYRAG	B.burgdorferi gyrA gene encoding DNA gyrase subunit A (partial)	96	289
2	97	100759	98837	gb|U04527|	Borrelia burgdorferi 212 DNA gyrase b subunit (gyrB) and ribonuclease P protein component (rnpA) genes, partial cds, DnaA protein (dnaA), DNA polymerase III beta subunit (dnaN), and ribosomal protein L34 (rpmH) genes, complete cds	98	904
2	98	100893	102389	gb|U04527|	Borrelia burgdorferi 212 DNA gyrase b subunit (gyrB) and ribonuclease P protein component (rnpA) genes, partial cds, DnaA protein (dnaA), DNA polymerase III beta subunit (dnaN), and ribosomal protein L34 (rpmH) genes, complete cds	100	1497
2	99	102618	103787	gb|U04527|	Borrelia burgdorferi 212 DNA gyrase b subunit (gyrB) and ribonuclease P protein component (rnpA) genes, partial cds, DnaA protein (dnaA), DNA polymerase III beta subunit (dnaN), and ribosomal protein L34 (rpmH) genes, complete cds	99	1170
2	100	103786	103607	gb|U04527|	Borrelia burgdorferi 212 DNA gyrase b subunit (gyrB) and ribonuclease P protein component (rnpA) genes, partial cds, DnaA protein (dnaA), DNA polymerase III beta subunit (dnaN), and ribosomal protein L34 (rpmH) genes, complete cds	100	180
2	101	103866	104177	gb|U04527|	Borrelia burgdorferi 212 DNA gyrase b subunit (gyrB) and ribonuclease P protein component (rnpA) genes, partial cds, DnaA protein (dnaA), DNA polymerase III beta subunit (dnaN), and	100	312

					ribosomal protein L34 (rpmH) genes, complete cds		
2	102	104254	104424	gb|U04527|	Borrelia burgdorferi 212 DNA gyrase b subunit (gyrB) and ribonuclease P protein component (rnpA) genes, partial cds, DnaA protein (dnaA), DNA polymerase III beta subunit (dnaN), and ribosomal protein L34 (rpmH) genes, complete cds	100	171
2	103	104393	104764	gb|U04527|	Borrelia burgdorferi 212 DNA gyrase b subunit (gyrB) and ribonuclease P protein component (rnpA) genes, partial cds, DnaA protein (dnaA), DNA polymerase III beta subunit (dnaN), and ribosomal protein L34 (rpmH) genes, complete cds	100	148
3	11	7200	8597	gb|U43739|	Borrelia burgdorferi fesmid clone 31, complete sequence	100	1185
3	12	8581	9666	emb|X96685|BBCDG	B.burgdorferi cell division genes	99	912
3	13	9664	10767	emb|X96685|BBCDG	B.burgdorferi cell division genes	99	1104
3	14	10826	10614	emb|X96433|BBFTSWQA	B.burgdorferi ftsW, ftsQ & ftsA genes	100	213
3	15	10797	11546	emb|X96433|BBFTSWQA	B.burgdorferi ftsW, ftsQ & ftsA genes	99	750
3	16	11519	12787	gb|U43739|	Borrelia burgdorferi fesmid clone 31, complete sequence	100	1269
3	17	12785	14008	gb|U43739|	Borrelia burgdorferi fesmid clone 31, complete sequence	100	1224
3	18	14006	14701	gb|U43739|	Borrelia burgdorferi fesmid clone 31, complete sequence	100	696
3	19	14699	15655	gb|L76303|	Borrelia burgdorferi ftsA gene, 3' end of cds, ftsZ, orf230, smf, hslVU, flgBCE, fliEFGHI, flbABC genes, complete cds	98	712
3	20	15653	16213	gb|U43739|	Borrelia burgdorferi fesmid clone 31, complete sequence	100	561
3	21	16149	17552	emb|X96685|BBCDG	B.burgdorferi cell division genes	97	1404

3	22	17550	17993	gb|U43739|	Borrelia burgdorferi fesmid clone 31, complete sequence	100	444
3	23	17996	18475	gb|U43739|	Borrelia burgdorferi fesmid clone 31, complete sequence	100	480
3	24	18445	18822	gb|L76303|	Borrelia burgdorferi ftsA gene, 3' end of cds, ftsZ, orf230, smf, hslVU, flgBCE, fliEFGHI, flbABC genes, complete cds	100	378
3	25	18777	20546	gb|L76303|	Borrelia burgdorferi ftsA gene, 3' end of cds, ftsZ, orf230, smf, hslVU, flgBCE, fliEFGHI, flbABC genes, complete cds	100	1770
3	26	20544	21596	gb|U43739|	Borrelia burgdorferi fesmid clone 31, complete sequence	100	1053
3	27	21575	22531	gb|U43739|	Borrelia burgdorferi fesmid clone 31, complete sequence	100	957
3	28	22529	23860	gb|L76303|	Borrelia burgdorferi ftsA gene, 3' end of cds, ftsZ, orf230, smf, hslVU, flgBCE, fliEFGHI, flbABC genes, complete cds	99	1332
3	29	23836	24288	gb|L76303|	Borrelia burgdorferi ftsA gene, 3' end of cds, ftsZ, orf230, smf, hslVU, flgBCE, fliEFGHI, flbABC genes, complete cds	99	453
3	30	24269	24898	gb|U43739|	Borrelia burgdorferi fesmid clone 31, complete sequence	100	630
3	31	24864	26084	gb|L76303|	Borrelia burgdorferi ftsA gene, 3' end of cds, ftsZ, orf230, smf, hslVU, flgBCE, fliEFGHI, flbABC genes, complete cds	99	1221
3	32	26095	26541	gb|L76303|	Borrelia burgdorferi ftsA gene, 3' end of cds, ftsZ, orf230, smf, hslVU, flgBCE, fliEFGHI, flbABC genes, complete cds	100	447
3	33	26525	27874	gb|U43739|	Borrelia burgdorferi fesmid clone 31, complete sequence	100	1350
3	34	27891	28121	gb|U43739|	Borrelia burgdorferi fesmid clone 31, complete sequence	100	231
3	35	28112	28900	gb|U43739|	Borrelia burgdorferi fesmid clone 31, complete sequence	100	789

3	36	28894	29682	gb|U43739|	Borrelia burgdorferi fesmid clone 31, complete sequence	100	789
3	37	29680	30267	gb|U43739|	Borrelia burgdorferi fesmid clone 31, complete sequence	100	588
3	38	30302	31363	gb|U43739|	Borrelia burgdorferi fesmid clone 31, complete sequence	100	1062
3	39	31403	31750	gb|U43739|	Borrelia burgdorferi fesmid clone 31, complete sequence	100	348
3	40	31728	32369	gb|U43739|	Borrelia burgdorferi fesmid clone 31, complete sequence	100	642
3	41	32356	33144	gb|L75945|	Borrelia burgdorferi flagellar hook protein (flgE), flbD, flagellar motor apparatus (motAB), fliL, fliM, fliZ, flagellar export apparatus (fliPQR, flhB), flhF, flbE genes	99	789
3	42	32642	32436	gb|L75945|	Borrelia burgdorferi flagellar hook protein (flgE), flbD, flagellar motor apparatus (motAB), fliL, fliM, fliZ, flagellar export apparatus (fliPQR, flhB), flhF, flbE genes	100	207
3	43	33129	33416	gb|L75945|	Borrelia burgdorferi flagellar hook protein (flgE), flbD, flagellar motor apparatus (motAB), fliL, fliM, fliZ, flagellar export apparatus (fliPQR, flhB), flhF, flbE genes	100	288
3	44	33425	34237	gb|L75945|	Borrelia burgdorferi flagellar hook protein (flgE), flbD, flagellar motor apparatus (motAB), fliL, fliM, fliZ, flagellar export apparatus (fliPQR, flhB), flhF, flbE genes	100	813
3	45	34320	34072	gb|L75945|	Borrelia burgdorferi flagellar hook protein (flgE), flbD, flagellar motor apparatus (motAB), fliL, fliM, fliZ, flagellar export apparatus (fliPQR, flhB), flhF, flbE genes	100	249
3	46	34222	35355	gb|L75945|	Borrelia burgdorferi flagellar hook protein (flgE), flbD, flagellar motor apparatus (motAB), fliL, fliM, fliZ, flagellar export apparatus (fliPQR, flhB), flhF, flbE genes	99	1134

Borrelia burgdorferi - Coding regions containing to know proteins

3	47	35349	37457	gb|L75945|	Borrelia burgdorferi flagellar hook protein (flgE), flbD, flagellar motor apparatus (motAB), fliL, fliM, fliZ, flagellar export apparatus (fliPQR, flhB), flhF, flbE genes	100	2109
3	48	37455	38627	gb|U43739|	Borrelia burgdorferi fesmid clone 31, complete sequence	100	1173
3	49	38612	39526	gb|U43739|	Borrelia burgdorferi fesmid clone 31, complete sequence	99	816
3	50	39765	39421	gb|U43739|	Borrelia burgdorferi fesmid clone 31, complete sequence	100	345
3	51	39524	40012	gb|U43739|	Borrelia burgdorferi fesmid clone 31, complete sequence	100	489
3	52	39985	41919	gb|U43739|	Borrelia burgdorferi fesmid clone 31, complete sequence	100	1935
3	53	41928	42248	gb|U43739|	Borrelia burgdorferi fesmid clone 31, complete sequence	100	286
3	58	47505	46090	gb|M28682|	B.burgdorferei promoter element DNA	100	78
3	68	57596	60034	gb|L77216|	Borrelia burgdorferi (strain B31) protease (lon) gene, complete cds	99	2439
3	69	62374	60023	gb|L77216|	Borrelia burgdorferi (strain B31) protease (lon) gene, complete cds	100	274
3	92	82181	83026	gb|U35673|	Borrelia burgdorferi OrfR gene, partial cds, and S20, Hbb, OrfH and Rho genes, complete cds	99	542
3	93	83032	83358	gb|U35673|	Borrelia burgdorferi OrfR gene, partial cds, and S20, Hbb, OrfH and Rho genes, complete cds	100	327
3	94	83365	83697	gb|U48651|	Borrelia burgdorferi P1G histone-like protein HBbu (hbb) gene, complete cds	100	327
3	95	83695	84105	gb|U35673|	Borrelia burgdorferi OrfR gene, partial cds, and S20, Hbb, OrfH and Rho genes, complete cds	99	411
3	96	84167	85732	gb|U35673|	Borrelia burgdorferi OrfR gene, partial cds, and S20, Hbb, OrfH and Rho genes, complete cds	99	1566
3	97	85778	86038	gb|U35673|	Borrelia burgdorferi OrfR gene, partial cds, and S20, Hbb, OrfH and Rho genes, complete cds	100	106

4	2	1935	1147	gb|U61142|	Borrelia burgdorferi outer membrane porin protein Oms28 precursor (oms28) gene, complete cds	99	789
4	12	10037	11002	gb|U59487|	Borrelia burgdorferi P35 antigen protein gene, and 7.5 kDa lipoprotein gene, complete cds	100	966
4	13	11365	11153	gb|U59859|	Borrelia burgdorferi strain B31 6.6 kDa lipoprotein gene, complete cds	100	213
4	14	11577	12230	gb|U59487|	Borrelia burgdorferi P35 antigen protein gene, and 7.5 kDa lipoprotein gene, complete cds	100	373
4	15	12578	13414	gb|M85216|	Borrelia burgdorferi 27kD protein antigen gene (p27), complete cds	78	370
4	16	13511	13753	gb|U22451|	Borrelia burgdorferi 49kb linear plasmid small 12kDa lipoprotein gene, complete cds	99	243
4	23	18668	17793	gb|L31427|	Borrelia burgdorferi (clone BbK2.1) phoA fusion protein gene, partial cds	100	169
4	49	36694	36347	gb|U75867|	Borrelia burgdorferi decorin binding protein B (DbpB) gene, complete cds	99	329
4	50	36351	36929	gb|U75867|	Borrelia burgdorferi decorin binding protein B (DbpB) gene, complete cds	99	564
4	51	36838	36692	gb|U75867|	Borrelia burgdorferi decorin binding protein B (DbpB) gene, complete cds	100	147
4	52	37001	37624	gb|U75866|	Borrelia burgdorferi decorin binding protein A (DbpA) gene, complete cds	93	533
4	55	40073	39318	gb|U42599|	Borrelia burgdorferi plasmid cp18, OspE (ospE) gene, partial cds	69	731
4	58	43349	42447	gb|L23137|	Borrelia burgdorferi (27985CT2) OspA gene, 3' end and OspB gene, complete cds	99	903
4	59	44228	43347	emb|A04009|A04009	B.burgdorferi OspA gene and 5'flanking region	100	882
4	60	44792	44403	gb|L19702|	Borrelia burgdorferi outer surface protein A (ospA) and outer surface protein B (ospB) genes, complete cds	88	370
4	61	45198	44758	gb|L19702|	Borrelia burgdorferi outer surface protein A (ospA) and outer surface protein B (ospB) genes, complete	89	375

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					cds		
4	62	46440	45382	gb|L19702|	Borrelia burgdorferi outer surface protein A (ospA) and outer surface protein B (ospB) genes, complete cds	85	622
4	67	49363	50622	gb|L34016|	Borrelia burgdorferi (clone 8) S1 gene, complete cds	99	1260
4	68	50708	51580	gb|L34017|	Borrelia burgdorferi (clone 8) S2 gene, complete cds	99	837
4	69	52203	51655	gb|L31423|	Borrelia burgdorferi (clone BbK2.14) phoA fusion protein gene, partial cds	99	292
4	70	53018	52488	gb|L41151|	Borrelia burgdorferi (clone 8) s3 gene, complete cds	99	297
5	1	535	71	gb|U60642|	Borrelia burgdorferi plasmid cp32-4, sequence at position 4-6kb	91	465
5	2	1526	546	gb|U60642|	Borrelia burgdorferi plasmid cp32-4, sequence at position 4-6kb	89	374
5	4	2395	2129	gb|L31425|	Borrelia burgdorferi (clone BbK3.168) phoA fusion protein gene, partial cds	98	135
5	11	6832	6542	gb|S66708|	{target sequence for detection of Lyme disease agent} [Borrelia burgdorferi, B31, 30-kb circular plasmid pIP87, Plasmid, 416 nt]	97	290
5	12	7422	6817	gb|U44914|	Borrelia burgdorferi plasmid cp32-2, erpC and erpD genes, complete cds	87	595
5	13	8167	7565	gb|U44914|	Borrelia burgdorferi plasmid cp32-2, erpC and erpD genes, complete cds	84	147
5	14	9408	8284	gb|U44914|	Borrelia burgdorferi plasmid cp32-2, erpC and erpD genes, complete cds	72	568
5	15	10122	9427	gb|U30617|	Borrelia burgdorferi Bbk2.11 (bbk2.10), complete cds	93	560
5	16	10533	11324	gb|U44912|	Borrelia burgdorferi plasmid cp32-1, erpA and erpB genes, complete cds	93	790
5	17	11590	11330	gb|U44913|	Borrelia burgdorferi plasmid cp32-4, erpH gene, complete cds	95	261

5	18	11761	11588	gb|U42599|	Borrelia burgdorferi plasmid cp18, OspE (ospE) gene, partial cds	96	173
5	19	13256	11808	gb|U42599|	Borrelia burgdorferi plasmid cp18, OspE (ospE) gene, partial cds	95	1431
5	20	14187	13636	emb|X87201|BBRGABCD	B.burgdorferi plasmid, orfA, B, C, D, E, & F genes, clone pOMB14 and pOMB17	100	552
5	21	14727	14185	emb|X87201|BBRGABCD	B.burgdorferi plasmid, orfA, B, C, D, E, & F genes, clone pOMB14 and pOMB17	100	511
5	22	15588	14788	emb|X87201|BBRGABCD	B.burgdorferi plasmid, orfA, B, C, D, E, & F genes, clone pOMB14 and pOMB17	99	801
5	23	16097	15519	emb|X87201|BBRGABCD	B.burgdorferi plasmid, orfA, B, C, D, E, & F genes, clone pOMB14 and pOMB17	98	579
5	24	17276	16158	emb|X87201|BBRGABCD	B.burgdorferi plasmid, orfA, B, C, D, E, & F genes, clone pOMB14 and pOMB17	94	1075
5	25	17558	18526	gb|U45425|	Borrelia burgdorferi 2.9-5 locus, ORF-A-D, REP+, REP-, and lipoprotein (LP) genes, complete cds	86	927
5	26	19040	18564	gb|U45422|	Borrelia burgdorferi 2.9-2 locus, ORF-C gene, partial cds, ORF-D, REP+, REP-, and lipoprotein (LP) genes, complete cds	89	379
5	27	19712	19116	gb|U45421|	Borrelia burgdorferi 2.9-1 locus, ORF 5-8, ORF-A-D, REP+, REP-, and lipoprotein (LP) genes, complete cds	85	596
5	28	20164	19775	gb|U45421|	Borrelia burgdorferi 2.9-1 locus, ORF 5-8, ORF-A-D, REP+, REP-, and lipoprotein (LP) genes, complete cds	97	390
5	29	20504	20121	gb|U45426|	Borrelia burgdorferi 2.9-6 locus, ORF-A-D genes, complete cds and REP+ gene, partial cds	99	384
5	30	20799	20446	gb|U96714|	Borrelia burgdorferi B31 BlyA (blyA) and BlyB (blyB) genes, complete cds	99	354
5	31	21006	20797	gb|U45426|	Borrelia burgdorferi 2.9-6 locus, ORF-A-D genes, complete cds and REP+ gene, partial cds	97	210
5	32	21903	21076	gb|U96714|	Borrelia burgdorferi B31 BlyA (blyA) and BlyB	95	440

Borrelia burgdorferi - Coding regions containing to know proteins

					(blyB) genes, complete cds		
5	33	21470	21625	gb|U96714|	Borrelia burgdorferi B31 BlyA (blyA) and BlyB (blyB) genes, complete cds	94	151
5	34	22518	22051	gb|U45421|	Borrelia burgdorferi 2.9-1 locus, ORF 5-8, ORF-A-D, REP+, REP-, and lipoprotein (LP) genes, complete cds	90	467
5	35	22806	22516	gb|U45421|	Borrelia burgdorferi 2.9-1 locus, ORF 5-8, ORF-A-D, REP+, REP-, and lipoprotein (LP) genes, complete cds	93	286
5	36	23082	22840	gb|U45421|	Borrelia burgdorferi 2.9-1 locus, ORF 5-8, ORF-A-D, REP+, REP-, and lipoprotein (LP) genes, complete cds	95	242
5	37	23397	23080	gb|U45421|	Borrelia burgdorferi 2.9-1 locus, ORF 5-8, ORF-A-D, REP+, REP-, and lipoprotein (LP) genes, complete cds	96	317
5	38	23768	23388	gb|U45421|	Borrelia burgdorferi 2.9-1 locus, ORF 5-8, ORF-A-D, REP+, REP-, and lipoprotein (LP) genes, complete cds	95	381
5	39	24331	23750	gb|U45421|	Borrelia burgdorferi 2.9-1 locus, ORF 5-8, ORF-A-D, REP+, REP-, and lipoprotein (LP) genes, complete cds	90	495
5	51	29986	29417	gb|U60642|	Borrelia burgdorferi plasmid cp32-4, sequence at position 4-6kb	97	300
5	52	30414	29980	gb|U60642|	Borrelia burgdorferi plasmid cp32-4, sequence at position 4-6kb	99	435
5	53	30803	30357	gb|U60642|	Borrelia burgdorferi plasmid cp32-4, sequence at position 4-6kb	97	447
5	54	31204	30740	gb|U60642|	Borrelia burgdorferi plasmid cp32-4, sequence at position 4-6kb	96	465
5	55	31775	31215	gb|U60642|	Borrelia burgdorferi plasmid cp32-4, sequence at position 4-6kb	98	374
5	59	33577	32804	gb|L31425|	Borrelia burgdorferi (clone BbK3.168) phoA fusion protein gene, partial cds	87	135

6	16	9678	9022	gb|U01894|	Borrelia burgdorferi B31 outer surface protein C (ospC) gene, complete cds	100	657
6	17	9836	11425	gb|L25883|	Borrelia burgdorferi 26 kb plasmid GMP synthetase (guaA) gene, complete cds	98	1590
6	18	11435	12664	gb|U13372|	Borrelia burgdorferi 26 kb plasmid IMP dehydrogenase (guaB) gene, partial cds	100	1212
6	19	12195	11686	gb|U13372|	Borrelia burgdorferi 26 kb plasmid IMP dehydrogenase (guaB) gene, partial cds	100	510
7	1	695	3	gb|U85588|	Borrelia burgdorferi transposase-like protein (tra) gene, partial cds	99	693
7	2	1081	677	gb|U85588|	Borrelia burgdorferi transposase-like protein (tra) gene, partial cds	98	375
7	39	25041	25847	gb|U45423|	Borrelia burgdorferi 2.9-3 locus, ORF-C gene, partial cds, ORF-D, REP+, REP-, and lipoprotein (LP) genes, complete cds	77	437
8	2	1420	746	gb|U45424|	Borrelia burgdorferi 2.9-4 locus, ORF-C gene, partial cds, ORF-D, REP+, REP-, and lipoprotein (LP) genes, complete cds	80	193
8	12	14287	14087	gb|U84396|	Borrelia burgdorferi 16 kb plasmid DNA fragment	95	140
8	17	18352	17876	gb|U85588|	Borrelia burgdorferi transposase-like protein (tra) gene, partial cds	96	362
9	1	2815	2507	gb|U43414|	Borrelia burgdorferi linear plasmid lp16 DNA, complete sequence	100	309
9	2	3522	2767	gb|U43414|	Borrelia burgdorferi linear plasmid lp16 DNA, complete sequence	99	756
9	3	5188	5862	gb|U43414|	Borrelia burgdorferi linear plasmid lp16 DNA, complete sequence	100	675
9	4	6809	7255	gb|U43414|	Borrelia burgdorferi linear plasmid lp16 DNA, complete sequence	100	447
9	5	8621	7467	gb|U43414|	Borrelia burgdorferi linear plasmid lp16 DNA, complete sequence	100	1155
9	6	9079	8735	gb|U43414|	Borrelia burgdorferi linear plasmid lp16 DNA, complete sequence	100	345

9	7	10224	9214	gb|U43414|	Borrelia burgdorferi linear plasmid lp16 DNA, complete sequence	100	911
9	8	10370	10972	gb|U43414|	Borrelia burgdorferi linear plasmid lp16 DNA, complete sequence	100	603
9	9	11844	11107	gb|U43414|	Borrelia burgdorferi linear plasmid lp16 DNA, complete sequence	100	738
9	10	13299	13027	gb|U43414|	Borrelia burgdorferi linear plasmid lp16 DNA, complete sequence	99	273
9	11	13612	13241	gb|U43414|	Borrelia burgdorferi linear plasmid lp16 DNA, complete sequence	99	372
10	2	2164	1604	gb|U45422|	Borrelia burgdorferi 2.9-2 locus, ORF-C gene, partial cds, ORF-D, REP+, REP-, and lipoprotein (LP) genes, complete cds	78	278
10	3	2686	2886	gb|U12332|	Borrelia burgdorferi 16 kb plasmid hypothetical protein gene, complete cds	97	143
13	1	3	842	gb|U03641|	Borrelia burgdorferi lp21 circular plasmid, complete sequence	91	290
13	2	1525	983	gb|L16625|	Borrelia burgdorferi exported neurotoxin-like protein gene, complete cds	99	531
13	6	4098	4901	gb|U03641|	Borrelia burgdorferi lp21 circular plasmid, complete sequence	86	713
13	7	4691	4467	gb|U03641|	Borrelia burgdorferi lp21 circular plasmid, complete sequence	88	224
13	8	6348	5041	gb|U42599|	Borrelia burgdorferi plasmid cp18, OspE (ospE) gene, partial cds	82	1202
13	9	6673	7788	emb|X87127|BBPBRGEA	B.burgdorferi repeated DNA element, 30.5 kb circular plasmid copy	81	519
13	10	7786	8355	gb|U03641|	Borrelia burgdorferi lp21 circular plasmid, complete sequence	78	414
13	11	8393	8968	gb|U03641|	Borrelia burgdorferi lp21 circular plasmid, complete sequence	84	576
13	12	9290	9544	gb|U03641|	Borrelia burgdorferi lp21 circular plasmid, complete sequence	91	210

Borrelia burgdorferi - Coding regions containing to know proteins

14	6	5768	6217	gb|L31616|	Borrelia burgdorferi protein p23 gene, complete cds	89	396
14	7	6126	6671	gb|L31616|	Borrelia burgdorferi protein p23 gene, complete cds	85	242
16	5	3660	2854	gb|M97452|	Borrelia burgdorferi outer surface protein D (ospD) gene, complete cds	100	807
19	3	3136	3657	gb|L41151|	Borrelia burgdorferi (clone 8) s3 gene, complete cds	77	267
21	1	849	4	gb|U60963|	Borrelia burgdorferi plasmid cp32-1 PCR target site, partial sequence	95	296
21	2	1427	834	gb|U44914|	Borrelia burgdorferi plasmid cp32-2, erpC and erpD genes, complete cds	93	594
21	3	2168	1581	gb|U44914|	Borrelia burgdorferi plasmid cp32-2, erpC and erpD genes, complete cds	83	130
21	4	2946	2257	gb|U80956|	Borrelia burgdorferi strain 297CH putative outer membrane protein (ospF) gene, complete cds	91	350
21	5	3794	2964	gb|U44913|	Borrelia burgdorferi plasmid cp32-4, erpH gene, complete cds	100	401
21	6	4334	5143	gb|U44913|	Borrelia burgdorferi plasmid cp32-4, erpH gene, complete cds	99	413
21	7	5362	5183	gb|U44913|	Borrelia burgdorferi plasmid cp32-4, erpH gene, complete cds	91	180
21	8	5581	5360	gb|U42599|	Borrelia burgdorferi plasmid cp18, OspE (ospE) gene, partial cds	97	221
22	1	306	4	gb|U45421|	Borrelia burgdorferi 2.9-1 locus, ORF 5-8, ORF-A-D, REP+, REP-, and lipoprotein (LP) genes, complete cds	95	303
22	2	664	317	gb|U45421|	Borrelia burgdorferi 2.9-1 locus, ORF 5-8, ORF-A-D, REP+, REP-, and lipoprotein (LP) genes, complete cds	97	336
22	3	1230	658	gb|U45421|	Borrelia burgdorferi 2.9-1 locus, ORF 5-8, ORF-A-D, REP+, REP-, and lipoprotein (LP) genes, complete cds	89	406
24	1	69	4058	gb|U76406|	Borrelia burgdorferi putative vls recombination cassettes Vls2-Vls16b (vls) gene, complete	97	2305

					sequence		
24	2	4056	5108	gb|U76406|	Borrelia burgdorferi putative vls recombination cassettes Vls2-Vls16b (vls) gene, complete sequence	94	750
25	1	383	760	gb|U43414|	Borrelia burgdorferi linear plasmid lp16 DNA, complete sequence	100	378
25	2	1333	1536	gb|U43414|	Borrelia burgdorferi linear plasmid lp16 DNA, complete sequence	100	204
26	1	684	82	emb|X87127|BBPBRGEA	B.burgdorferi repeated DNA element, 30.5 kb circular plasmid copy	96	603
26	2	903	682	emb|X87127|BBPBRGEA	B.burgdorferi repeated DNA element, 30.5 kb circular plasmid copy	96	221
26	4	2181	2573	gb|AF000270|	Borrelia burgdorferi strain B31 2.9-like locus, OrfC, OrfD, Rev (rev), lipoprotein (LP), and 36 kDa-like orf2 genes, complete cds, and 36 kDa-like orf1 gene, partial cds	94	362
26	5	3073	2621	gb|U45427|	Borrelia burgdorferi 2.9-7 locus, ORF-A-D, REV, and lipoprotein (LPA and LPB) genes, complete cds	80	220
26	6	3745	3149	gb|U45423|	Borrelia burgdorferi 2.9-3 locus, ORF-C gene, partial cds, ORF-D, REP+, REP-, and lipoprotein (LP) genes, complete cds	87	478
26	8	4663	4355	gb|U45424|	Borrelia burgdorferi 2.9-4 locus, ORF-C gene, partial cds, ORF-D, REP+, REP-, and lipoprotein (LP) genes, complete cds	98	309
27	1	997	434	gb|L31615|	Borrelia burgdorferi (clone BbK2.5-6) unknown protein gene, complete cds	96	219
27	2	1395	2258	gb|L31616|	Borrelia burgdorferi protein p23 gene, complete cds	98	610
30	1	252	686	emb|X87127|BBPBRGEA	B.burgdorferi repeated DNA element, 30.5 kb circular plasmid copy	97	419
30	2	760	1545	emb|X87127|BBPBRGEA	B.burgdorferi repeated DNA element, 30.5 kb circular plasmid copy	100	786
30	3	1543	2157	emb|X87127|B	B.burgdorferi repeated DNA element, 30.5 kb	100	615

				BPBRGEA	circular plasmid copy		
30	4	2158	2802	emblX87127|BBPBRGEA	B.burgdorferi repeated DNA element, 30.5 kb circular plasmid copy	100	645
30	5	3247	4230	gb|U42599|	Borrelia burgdorferi plasmid cp18, OspE (ospE) gene, partial cds	95	976
33	1	450	995	gb|U72996|	Borrelia burgdorferi plasmid cp32-5, erpI gene, complete cds	100	546
33	2	1008	2159	gb|U78764|	Borrelia burgdorferi plasmid cp32-1, erpA and erpB2 genes, complete cds	100	1152
33	3	2253	2882	gb|U44914|	Borrelia burgdorferi plasmid cp32-2, erpC and erpD genes, complete cds	98	379
33	4	3050	3628	gb|U44914|	Borrelia burgdorferi plasmid cp32-2, erpC and erpD genes, complete cds	93	577
35	1	3	176	gb|U03396|	Borrelia burgdorferi B31 Ala-tRNA (alaT), Ile-tRNA (ileT), 16S rRNA, 23S rRNA (rrlA and rrlB), and 5S rRNA (rrfA and rrfB) genes, complete sequence	91	174
35	2	976	737	gb|M88330|	Borrelia burgdorferi 23S ribosomal RNA gene	100	240
36	1	1	525	emblX87201|BBRGABCD	B.burgdorferi plasmid, orfA, B, C, D, E, & F genes, clone pOMB14 and pOMB17	77	159
38	1	672	28	gb|U44914|	Borrelia burgdorferi plasmid cp32-2, erpC and erpD genes, complete cds	92	571
38	2	850	653	gb|U42598|	Borrelia burgdorferi plasmid cp32-3, ErpG (erpG) and BapA (bapA) genes, complete cds	100	133
38	3	1516	983	emblX82409|BBOSPG	B.burgdorferi ospG and bapA genes	100	534
38	4	2200	1604	emblX82409|BBOSPG	B.burgdorferi ospG and bapA genes	100	597
38	5	2602	3132	gb|U42598|	Borrelia burgdorferi plasmid cp32-3, ErpG (erpG) and BapA (bapA) genes, complete cds	99	529
39	1	967	665	emblX87202|BBBRGBCDE	B.burgdorferi plasmid, orfA, B, C, D, E, & G genes, clone pOMB10	97	170
39	2	1505	957	emblX87202|B	B.burgdorferi plasmid, orfA, B, C, D, E, & G	89	176

				BBRGBCDE	genes, clone pOMB10		
39	3	2353	1553	gb|U42599|	Borrelia burgdorferi plasmid cp18, OspE (ospE) gene, partial cds	91	137
39	4	2574	2284	emb|X87201|BBBRGABCD	B.burgdorferi plasmid, orfA, B, C, D, E, & F genes, clone pOMB14 and pOMB17	94	291
39	5	2874	2572	emb|X87127|BBPBRGEA	B.burgdorferi repeated DNA element, 30.5 kb circular plasmid copy	91	284
39	6	3028	2861	emb|X87127|BBPBRGEA	B.burgdorferi repeated DNA element, 30.5 kb circular plasmid copy	93	168
40	1	596	132	gb|U76406|	Borrelia burgdorferi putative vls recombination cassettes Vls2-Vls16b (vls) gene, complete sequence	100	465
40	2	1753	575	gb|U76406|	Borrelia burgdorferi putative vls recombination cassettes Vls2-Vls16b (vls) gene, complete sequence	100	1179
40	3	3000	1732	gb|U76406|	Borrelia burgdorferi putative vls recombination cassettes Vls2-Vls16b (vls) gene, complete sequence	99	1269
41	1	1	411	emb|X87201|BBBRGABCD	B.burgdorferi plasmid, orfA, B, C, D, E, & F genes, clone pOMB14 and pOMB17	99	411
41	2	342	1127	emb|X87201|BBBRGABCD	B.burgdorferi plasmid, orfA, B, C, D, E, & F genes, clone pOMB14 and pOMB17	80	785
41	3	1172	1747	emb|X87201|BBBRGABCD	B.burgdorferi plasmid, orfA, B, C, D, E, & F genes, clone pOMB14 and pOMB17	80	572
41	4	1745	2338	gb|U42599|	Borrelia burgdorferi plasmid cp18, OspE (ospE) gene, partial cds	65	571
42	2	1133	1384	gb|U85588|	Borrelia burgdorferi transposase-like protein (tra) gene, partial cds	93	236
43	1	360	4	emb|X87127|BBPBRGEA	B.burgdorferi repeated DNA element, 30.5 kb circular plasmid copy	96	356
43	2	635	1741	gb|AF000270|	Borrelia burgdorferi strain B31 2.9-like locus, OrfC, OrfD, Rev (rev), lipoprotein (LP), and 36 kDa-like orf2 genes, complete cds, and 36 kDa-like	90	392

					orf1 gene, partial cds		
43	3	2242	1784	gb|U45423|	Borrelia burgdorferi 2.9-3 locus, ORF-C gene, partial cds, ORF-D, REP+, REP-, and lipoprotein (LP) genes, complete cds	85	421
43	4	2860	2318	gb|U45421|	Borrelia burgdorferi 2.9-1 locus, ORF 5-8, ORF-A-D, REP+, REP-, and lipoprotein (LP) genes, complete cds	95	259
44	1	1158	178	gb|U60642|	Borrelia burgdorferi plasmid cp32-4, sequence at position 4-6kb	89	374
44	3	2531	1761	gb|L31425|	Borrelia burgdorferi (clone BbK3.168) phoA fusion protein gene, partial cds	99	135
45	1	287	3	gb|U78764|	Borrelia burgdorferi plasmid cp32-1, erpA and erpB2 genes, complete cds	84	153
45	3	2037	1453	gb|L13924|	Borrelia burgdorferi outer surface protein E (OspE) gene, complete cds	90	386
45	4	2663	2893	gb|U44912|	Borrelia burgdorferi plasmid cp32-1, erpA and erpB genes, complete cds	90	230
46	1	174	338	gb|U42599|	Borrelia burgdorferi plasmid cp18, OspE (ospE) gene, partial cds	96	91
46	2	259	966	gb|U42599|	Borrelia burgdorferi plasmid cp18, OspE (ospE) gene, partial cds	100	692
46	3	964	1527	gb|U42599|	Borrelia burgdorferi plasmid cp18, OspE (ospE) gene, partial cds	100	564
46	4	1509	2111	gb|U42599|	Borrelia burgdorferi plasmid cp18, OspE (ospE) gene, partial cds	99	603
46	5	2537	2851	emb|X87201|BBRGABCD	B.burgdorferi plasmid, orfA, B, C, D, E, & F genes, clone pOMB14 and pOMB17	98	315
47	1	2	526	gb|U45425|	Borrelia burgdorferi 2.9-5 locus, ORF-A-D, REP+, REP-, and lipoprotein (LP) genes, complete cds	95	525
47	2	1245	724	gb|U45424|	Borrelia burgdorferi 2.9-4 locus, ORF-C gene, partial cds, ORF-D, REP+, REP-, and lipoprotein (LP) genes, complete cds	94	483

47	3	1971	1321	gb|U45424|	Borrelia burgdorferi 2.9-4 locus, ORF-C gene, partial cds, ORF-D, REP+, REP-, and lipoprotein (LP) genes, complete cds	89	651
48	1	363	25	gb|U44914|	Borrelia burgdorferi plasmid cp32-2, erpC and erpD genes, complete cds	87	327
48	2	412	1182	gb|U72997|	Borrelia burgdorferi plasmid cp32-6, erpK gene, complete cds	100	91
48	3	2047	1244	gb|U72997|	Borrelia burgdorferi plasmid cp32-6, erpK gene, complete cds	99	804
49	1	713	18	gb|U76406|	Borrelia burgdorferi putative vls recombination cassettes Vls2-Vls16b (vls) gene, complete sequence	99	606
49	2	2308	704	gb|U76406|	Borrelia burgdorferi putative vls recombination cassettes Vls2-Vls16b (vls) gene, complete sequence	98	1596
51	1	613	2	gb|U42599|	Borrelia burgdorferi plasmid cp18, OspE (ospE) gene, partial cds	99	612
51	3	2203	2487	gb|U44914|	Borrelia burgdorferi plasmid cp32-2, erpC and erpD genes, complete cds	86	269
52	1	3	236	emb|X87202|BBRGBCDE	B.burgdorferi plasmid, orfA, B, C, D, E, & G genes, clone pOMB10	94	146
52	2	179	319	emb|X87201|BBRGABCD	B.burgdorferi plasmid, orfA, B, C, D, E, & F genes, clone pOMB14 and pOMB17	94	140
52	3	250	1050	gb|U42599|	Borrelia burgdorferi plasmid cp18, OspE (ospE) gene, partial cds	86	146
52	6	1650	2201	gb|U42599|	Borrelia burgdorferi plasmid cp18, OspE (ospE) gene, partial cds	81	422
53	1	93	581	gb|AF000270|	Borrelia burgdorferi strain B31 2.9-like locus, OrfC, OrfD, Rev (rev), lipoprotein (LP), and 36 kDa-like orf2 genes, complete cds, and 36 kDa-like orf1 gene, partial cds	99	489
53	2	883	719	gb|AF000270|	Borrelia burgdorferi strain B31 2.9-like locus, OrfC, OrfD, Rev (rev), lipoprotein (LP), and 36 kDa-like orf2 genes, complete cds, and 36 kDa-like	100	101

Borrelia burgdorferi - Coding regions containing to know proteins

					orf1 gene, partial cds		
53	3	1107	811	gb|AF000270|	Borrelia burgdorferi strain B31 2.9-like locus, OrfC, OrfD, Rev (rev), lipoprotein (LP), and 36 kDa-like orf2 genes, complete cds, and 36 kDa-like orf1 gene, partial cds	100	289
53	4	1447	1064	gb|AF000270|	Borrelia burgdorferi strain B31 2.9-like locus, OrfC, OrfD, Rev (rev), lipoprotein (LP), and 36 kDa-like orf2 genes, complete cds, and 36 kDa-like orf1 gene, partial cds	96	381
53	5	1742	1380	gb|U45427|	Borrelia burgdorferi 2.9-7 locus, ORF-A-D, REV, and lipoprotein (LPA and LPB) genes, complete cds	93	362
53	6	1949	1740	gb|U45426|	Borrelia burgdorferi 2.9-6 locus, ORF-A-D genes, complete cds and REP+ gene, partial cds	98	210
57	1	3	434	gb|U45422|	Borrelia burgdorferi 2.9-2 locus, ORF-C gene, partial cds, ORF-D, REP+, REP-, and lipoprotein (LP) genes, complete cds	92	326
57	2	1580	471	gb|AF000270|	Borrelia burgdorferi strain B31 2.9-like locus, OrfC, OrfD, Rev (rev), lipoprotein (LP), and 36 kDa-like orf2 genes, complete cds, and 36 kDa-like orf1 gene, partial cds	98	362
57	3	1837	2109	emb|X87127|BBPBRGEA	B.burgdorferi repeated DNA element, 30.5 kb circular plasmid copy	84	246
58	4	1573	1800	gb|L31425|	Borrelia burgdorferi (clone BbK3.168) phoA fusion protein gene, partial cds	90	118
60	1	668	111	emb|X87201|BBBRGABCD	B.burgdorferi plasmid, orfA, B, C, D, E, & F genes, clone pOMB14 and pOMB17	75	519
60	2	1479	694	emb|X87127|BBPBRGEA	B.burgdorferi repeated DNA element, 30.5 kb circular plasmid copy	72	786
60	3	1907	1410	emb|X87201|BBBRGABCD	B.burgdorferi plasmid, orfA, B, C, D, E, & F genes, clone pOMB14 and pOMB17	95	498
62	1	284	3	emb|X87201|BBBRGABCD	B.burgdorferi plasmid, orfA, B, C, D, E, & F genes, clone pOMB14 and pOMB17	79	260
62	2	878	282	emb|X87202|B	B.burgdorferi plasmid, orfA, B, C, D, E, & G	74	501

Borrelia burgdorferi - Coding regions containing to know proteins

				BBRGBCDE	genes, clone pOMB10		
62	3	1704	910	gb|U42599|	Borrelia burgdorferi plasmid cp18, OspE (ospE) gene, partial cds	78	351
64	1	563	54	gb|U43414|	Borrelia burgdorferi linear plasmid lp16 DNA, complete sequence	100	510
64	2	1320	1117	gb|U43414|	Borrelia burgdorferi linear plasmid lp16 DNA, complete sequence	100	204
66	1	647	75	gb|U60642|	Borrelia burgdorferi plasmid cp32-4, sequence at position 4-6kb	93	300
66	2	1075	641	gb|U60642|	Borrelia burgdorferi plasmid cp32-4, sequence at position 4-6kb	96	435
66	3	1530	1018	gb|U60642|	Borrelia burgdorferi plasmid cp32-4, sequence at position 4-6kb	94	440
70	1	3	275	gb|U96714|	Borrelia burgdorferi B31 BlyA (blyA) and BlyB (blyB) genes, complete cds	98	207
70	2	217	600	gb|U45426|	Borrelia burgdorferi 2.9-6 locus, ORF-A-D genes, complete cds and REP+ gene, partial cds	99	384
70	3	557	946	gb|U45423|	Borrelia burgdorferi 2.9-3 locus, ORF-C gene, partial cds, ORF-D, REP+, REP-, and lipoprotein (LP) genes, complete cds	98	390
70	4	1424	1083	gb|AF000270|	Borrelia burgdorferi strain B31 2.9-like locus, OrfC, OrfD, Rev (rev), lipoprotein (LP), and 36 kDa-like orf2 genes, complete cds, and 36 kDa-like orf1 gene, partial cds	99	342
75	1	2	925	gb|U60642|	Borrelia burgdorferi plasmid cp32-4, sequence at position 4-6kb	98	374
75	2	936	1328	gb|U60642|	Borrelia burgdorferi plasmid cp32-4, sequence at position 4-6kb	96	393
76	1	464	12	gb|U45422|	Borrelia burgdorferi 2.9-2 locus, ORF-C gene, partial cds, ORF-D, REP+, REP-, and lipoprotein (LP) genes, complete cds	85	281
76	2	1256	540	gb|U45425|	Borrelia burgdorferi 2.9-5 locus, ORF-A-D, REP+, REP-, and lipoprotein (LP) genes, complete	91	552

					cds		
77	1	433	2	gb|U45422|	Borrelia burgdorferi 2.9-2 locus, ORF-C gene, partial cds, ORF-D, REP+, REP-, and lipoprotein (LP) genes, complete cds	90	379
77	2	1159	509	gb|U45424|	Borrelia burgdorferi 2.9-4 locus, ORF-C gene, partial cds, ORF-D, REP+, REP-, and lipoprotein (LP) genes, complete cds	97	651
81	2	657	1034	gb|U43414|	Borrelia burgdorferi linear plasmid lp16 DNA, complete sequence	80	255
83	1	3	1202	gb|U76406|	Borrelia burgdorferi putative vls recombination cassettes Vls2-Vls16b (vls) gene, complete sequence	99	1198
85	1	1	360	gb|U45421|	Borrelia burgdorferi 2.9-1 locus, ORF 5-8, ORF-A-D, REP+, REP-, and lipoprotein (LP) genes, complete cds	91	347
85	2	358	1008	gb|U96714|	Borrelia burgdorferi B31 BlyA (blyA) and BlyB (blyB) genes, complete cds	84	440
85	3	791	636	gb|U96714|	Borrelia burgdorferi B31 BlyA (blyA) and BlyB (blyB) genes, complete cds	80	151
86	1	891	289	gb|U45422|	Borrelia burgdorferi 2.9-2 locus, ORF-C gene, partial cds, ORF-D, REP+, REP-, and lipoprotein (LP) genes, complete cds	86	486
86	2	1151	954	gb|U45427|	Borrelia burgdorferi 2.9-7 locus, ORF-A-D, REV, and lipoprotein (LPA and LPB) genes, complete cds	97	148
88	1	137	3	gb|U60642|	Borrelia burgdorferi plasmid cp32-4, sequence at position 4-6kb	97	135
88	2	325	131	gb|U60642|	Borrelia burgdorferi plasmid cp32-4, sequence at position 4-6kb	98	195
88	3	565	323	gb|U60642|	Borrelia burgdorferi plasmid cp32-4, sequence at position 4-6kb	98	243
88	4	954	508	gb|U60642|	Borrelia burgdorferi plasmid cp32-4, sequence at position 4-6kb	98	447

Borrelia burgdorferi - Coding regions containing to know proteins

88	5	1091	891	gb|U60640|	Borrelia burgdorferi plasmid cp32-2, sequence at position 5kb	98	201
91	1	927	34	gb|U45422|	Borrelia burgdorferi 2.9-2 locus, ORF-C gene, partial cds, ORF-D, REP+, REP-, and lipoprotein (LP) genes, complete cds	97	313
93	1	162	578	gb|U45421|	Borrelia burgdorferi 2.9-1 locus, ORF 5-8, ORF-A-D, REP+, REP-, and lipoprotein (LP) genes, complete cds	89	331
93	2	572	940	gb|U45421|	Borrelia burgdorferi 2.9-1 locus, ORF 5-8, ORF-A-D, REP+, REP-, and lipoprotein (LP) genes, complete cds	96	368
94	1	3	245	gb|U45425|	Borrelia burgdorferi 2.9-5 locus, ORF-A-D, REP+, REP-, and lipoprotein (LP) genes, complete cds	97	243
94	2	749	282	gb|AF000270|	Borrelia burgdorferi strain B31 2.9-like locus, OrfC, OrfD, Rev (rev), lipoprotein (LP), and 36 kDa-like orf2 genes, complete cds, and 36 kDa-like orf1 gene, partial cds	90	458
97	1	506	3	gb|U44914|	Borrelia burgdorferi plasmid cp32-2, erpC and erpD genes, complete cds	94	472
98	1	827	264	gb|U42599|	Borrelia burgdorferi plasmid cp18, OspE (ospE) gene, partial cds	70	380
99	1	175	408	gb|U43414|	Borrelia burgdorferi linear plasmid lp16 DNA, complete sequence	100	234
99	2	329	757	gb|U43414|	Borrelia burgdorferi linear plasmid lp16 DNA, complete sequence	98	220
101	1	207	440	gb|U43414|	Borrelia burgdorferi linear plasmid lp16 DNA, complete sequence	100	234
101	2	361	837	gb|U43414|	Borrelia burgdorferi linear plasmid lp16 DNA, complete sequence	99	477
102	1	3	911	gb|U76406|	Borrelia burgdorferi putative vls recombination cassettes Vls2-Vls16b (vls) gene, complete sequence	99	889
104	1	388	242	gb|U45426|	Borrelia burgdorferi 2.9-6 locus, ORF-A-D genes,	100	146

Borrelia burgdorferi - Coding regions containing to know proteins

					complete cds and REP+ gene, partial cds		
104	2	595	386	gb|U96714|	Borrelia burgdorferi B31 BlyA (blyA) and BlyB (blyB) genes, complete cds	100	210
107	1	2	811	gb|U45425|	Borrelia burgdorferi 2.9-5 locus, ORF-A-D, REP+, REP-, and lipoprotein (LP) genes, complete cds	95	789
109	1	264	4	gb|L31616|	Borrelia burgdorferi protein p23 gene, complete cds	86	201
109	2	598	173	gb|L31616|	Borrelia burgdorferi protein p23 gene, complete cds	93	396
109	3	807	580	gb|L31615|	Borrelia burgdorferi (clone BbK2.5-6) unknown protein gene, complete cds	99	228
110	1	1	456	gb|U45421|	Borrelia burgdorferi 2.9-1 locus, ORF 5-8, ORF-A-D, REP+, REP-, and lipoprotein (LP) genes, complete cds	95	456
110	2	450	761	gb|U45421|	Borrelia burgdorferi 2.9-1 locus, ORF 5-8, ORF-A-D, REP+, REP-, and lipoprotein (LP) genes, complete cds	93	310
111	1	787	215	gb|U45421|	Borrelia burgdorferi 2.9-1 locus, ORF 5-8, ORF-A-D, REP+, REP-, and lipoprotein (LP) genes, complete cds	89	405
119	1	653	84	gb|U60642|	Borrelia burgdorferi plasmid cp32-4, sequence at position 4-6kb	98	300
121	1	719	123	gb|U60642|	Borrelia burgdorferi plasmid cp32-4, sequence at position 4-6kb	98	374
122	1	403	2	gb|U44914|	Borrelia burgdorferi plasmid cp32-2, erpC and erpD genes, complete cds	85	391
128	1	175	408	gb|U43414|	Borrelia burgdorferi linear plasmid lp16 DNA, complete sequence	100	234
128	2	329	700	gb|U43414|	Borrelia burgdorferi linear plasmid lp16 DNA, complete sequence	99	356
129	1	458	697	emb|X87201|BBBRGABCD	B.burgdorferi plasmid, orfA, B, C, D, E, & F genes, clone pOMB14 and pOMB17	100	238
132	1	234	467	gb|U43414|	Borrelia burgdorferi linear plasmid lp16 DNA, complete sequence	100	234

132	2	388	660	gb|U43414|	Borrelia burgdorferi linear plasmid lp16 DNA, complete sequence	99	171
133	1	3	560	emb|X87127|BBPBRGEA	B.burgdorferi repeated DNA element, 30.5 kb circular plasmid copy	80	243
134	1	339	4	emb|X87202|BBBRGBCDE	B.burgdorferi plasmid, orfA, B, C, D, E, & G genes, clone pOMB10	78	331
141	1	554	33	gb|U96714|	Borrelia burgdorferi B31 BlyA (blyA) and BlyB (blyB) genes, complete cds	100	513
141	2	124	276	gb|U96714|	Borrelia burgdorferi B31 BlyA (blyA) and BlyB (blyB) genes, complete cds	100	153
143	1	67	498	gb|U42598|	Borrelia burgdorferi plasmid cp32-3, ErpG (erpG) and BapA (bapA) genes, complete cds	98	432
144	1	497	3	emb|X87127|BBPBRGEA	B.burgdorferi repeated DNA element, 30.5 kb circular plasmid copy	94	495
146	1	193	2	gb|U03641|	Borrelia burgdorferi Ip21 circular plasmid, complete sequence	86	144
147	1	3	542	gb|U60642|	Borrelia burgdorferi plasmid cp32-4, sequence at position 4-6kb	88	296
153	1	352	2	gb|M96847|	Borrelia burgdorferi GrpE protein homologue gene, DnaK protein homologue gene, and DnaJ protein homologue gene, complete cds's	97	351

TABLE 6.

Borrelia burgdorferi - Putative coding regions of novel proteins not similar to know proteins

Contig ID	ORF ID	Start (nt)	Stop (nt)
2	4	2730	3554
2	5	3559	3410
2	7	5464	3869
2	13	10502	9999
2	17	13800	13576
2	19	15368	15204
2	28	21155	21400
2	50	41944	42186
2	58	53786	52911
2	59	54816	53773
2	61	57393	55813
2	63	57882	57682
2	65	60898	60203
2	66	61441	62070
2	67	62078	62692
2	70	65896	66540
2	74	70203	69910
2	78	71818	71399
2	80	72956	74032
2	81	73515	73267
2	90	92181	92525
2	91	92968	92555
2	108	109872	110057
2	112	112408	112812
2	113	112858	113037
2	114	113035	113460
2	115	113506	113724
2	119	114325	114852
3	6	3279	4079
3	8	5156	6019
3	54	42256	42789
3	59	47264	47506
3	60	47673	48692
3	63	51475	51026
3	70	60330	60575
3	71	61050	61349
3	72	61347	61670
3	74	63917	64303
3	86	75347	75532
3	88	76593	77384
3	99	89769	89005
3	102	91278	91661
3	103	92137	92463
3	105	92423	92785
3	108	93467	93886
3	115	98262	98681

Borrelia burgdorferi - Putative coding regions of novel proteins not similar to know proteins

3	121	102227	102904
3	126	111308	110055
4	6	3751	4179
4	7	4218	5042
4	19	16115	15516
4	20	17028	16075
4	21	17379	17092
4	22	17735	17397
4	24	19243	18785
4	25	18942	19196
4	26	20677	19259
4	27	19431	19751
4	29	21376	20876
4	30	21899	21423
4	31	22918	21845
4	33	23951	23553
4	37	26253	25627
4	38	26991	26332
4	39	28181	26931
4	40	29175	28522
4	43	30605	30342
4	45	34906	33548
4	48	35750	35932
5	3	2102	1527
5	5	2656	2393
5	7	3460	2900
5	10	6544	5645
5	40	25278	24322
5	41	25235	25600
5	42	25665	25276
5	44	25881	25663
5	47	27883	27410
5	48	28351	27881
5	49	29028	28324
5	50	29454	29026
5	56	32199	31666
5	57	32571	32200
5	58	32826	32569
5	60	32913	33245
5	61	33766	33575
5	62	34173	33742
5	64	35514	34861
6	2	954	1181
6	3	1590	1763
6	5	3400	3954
6	7	4691	5218
6	8	5187	5699

Borrelia burgdorferi - Putative coding regions of novel proteins not similar to know proteins

6	11	6498	5983
6	12	6975	6727
6	14	7978	7448
6	15	8479	7976
6	22	15106	15636
6	27	19999	18842
6	28	20036	20668
6	29	21814	20690
6	30	20949	21269
6	35	24136	23630
6	37	25697	26248
7	8	8100	7792
7	10	8145	8288
7	11	9374	8517
7	12	9771	9325
7	13	9652	10185
7	14	10163	9765
7	15	10517	10173
7	16	11363	10524
7	17	11904	11392
7	18	12495	11902
7	19	13516	12473
7	20	12807	13154
7	22	15149	14697
7	24	15855	15046
7	25	15503	15826
7	26	16638	15853
7	27	19344	16636
7	31	19473	19727
7	32	20067	19675
7	33	20762	20049
7	34	21136	20738
7	36	22975	23406
7	40	26667	25870
8	3	2907	4118
8	5	5898	6059
8	6	7399	8313
8	13	15645	15899
8	14	17281	16331
8	15	16905	17111
10	4	3211	3684
10	6	3857	4456
10	8	5982	5599
10	11	8038	7802
10	14	10255	10100
11	7	5688	5828
11	9	7248	7685

Borrelia burgdorferi - Putative coding regions of novel proteins not similar to know proteins

11	10	7672	8028
11	13	9642	10154
12	1	101	370
12	2	982	680
12	3	1390	1115
12	4	1528	1388
12	5	1913	1431
12	11	7308	6616
14	2	3588	3328
14	4	4657	4815
14	9	7981	8511
15	1	1	327
15	2	325	1077
15	3	1478	657
15	4	2360	1758
15	5	2839	2507
15	9	3922	3743
15	10	4145	3900
15	11	4112	4270
15	13	7677	6127
15	14	7852	7709
15	15	8052	7825
15	16	8222	7857
16	2	1733	1936
16	3	1905	2063
16	6	5212	4220
16	7	8903	8505
17	2	1500	1709
17	5	4097	4660
17	7	6344	6189
18	1	1635	2465
18	2	2509	3306
18	3	3332	4390
18	5	4933	4727
18	7	6353	7084
18	8	7098	7625
20	7	4700	4557
22	4	2175	1228
22	5	2132	2314
22	6	2829	2173
22	8	3254	3601
22	9	4408	4169
22	10	4875	4402
22	11	5343	4873
23	2	2283	1537
23	3	3564	2617
25	6	3677	4147

Borrelia burgdorferi - Putative coding regions of novel proteins not similar to know proteins

26	7	4251	3889
28	2	732	1739
29	3	310	885
31	1	28	195
32	3	935	1603
32	4	1637	2332
37	2	1379	1059
42	4	2708	2388
44	2	1734	1159
44	4	2942	2532
47	4	2336	2115
50	1	908	120
52	4	674	501
56	1	152	1465
56	2	611	459
56	3	1479	2150
58	3	1691	1329
58	5	1867	2046
59	2	2018	1044
61	1	1	657
61	3	1389	1907
62	4	1115	1345
63	1	663	325
63	2	769	446
63	3	1759	1013
65	1	472	903
65	2	901	1236
67	1	387	4
67	2	979	401
67	3	1482	961
68	2	451	612
69	3	840	574
71	1	363	4
72	1	586	933
73	1	300	4
73	2	824	279
73	3	1396	1145
79	1	22	1119
82	1	701	303
82	2	1188	775
84	1	331	134
84	2	983	348
87	1	277	2
87	2	1136	267
96	1	434	57
96	2	748	557
97	2	976	659

Borrelia burgdorferi - Putative coding regions of novel proteins not similar to know proteins

103	1	301	2
103	2	886	299
105	1	36	509
106	1	425	3
106	3	761	600
112	1	416	799
113	1	685	59
118	1	1	489
118	2	487	753
120	2	299	691
124	1	1	630
127	1	702	322
135	1	287	3
135	2	649	407
136	1	1	645
140	2	619	332
145	1	1	480

(1) GENERAL INFORMATION:

(i) APPLICANT: Human Genome Sciences, Inc. et al.

(ii) TITLE OF INVENTION: Borrelia burgdorferi Polynucleotides and Sequences

(iii) NUMBER OF SEQUENCES: 155

(iv) CORRESPONDENCE ADDRESS:

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(C) CITY: Rockville

(D) STATE: Maryland

(E) COUNTRY: USA

(F) ZIP: 20850

(v) COMPUTER READABLE FORM:

(A) MEDIUM TYPE: Diskette, 3.50 inch, 1.4Mb storage

(B) COMPUTER: HP Vectra 486/33

(C) OPERATING SYSTEM: MSDOS version 6.2

(D) SOFTWARE: ASCII Text

(vi) CURRENT APPLICATION DATA:

(A) APPLICATION NUMBER:

(B) FILING DATE: Herewith

(C) CLASSIFICATION:

(vii) PRIOR APPLICATION DATA:

(A) APPLICATION NUMBER:

(B) FILING DATE:

(viii) ATTORNEY/AGENT INFORMATION:

(A) NAME: Brookes, A. Anders

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(2) INFORMATION FOR SEQ ID NO: 1:

(i) SEQUENCE CHARACTERISTICS:
(A) LENGTH: 910715 base pairs
(B) TYPE: nucleic acid
(C) STRANDEDNESS: double
(D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 1:

ATATAATTTT TAATTAGTAT AGAATATGTT AAACTTTACC CTTGAATTTT TCTACTCTAT 60

TTGTATATTC TATAGAAAAA ACGATTAGAA TTAAACAAAG CCATAACTGA ACCAACGGTA 120

ATTAGTAGAT AAAGGGATCA AAATATTTTT TATTGCAGCA AGAATACCTT GGTATATTAG 180

AAAAACCAAA AGTCATAGTC AAATCATCTT TTGATAACAA TCCCCAAATC TATAATTTAT 240

TATGAAATTA ATTGCTCCCT TGAAAAGATT AGTTTTTAAA ACTACAAGAC TACTATCAAT 300

CACTATCAGA TAGATTAAAA CAACCTTTAC AAGAAAAAAA TCTTACTACT ATTTTATTGT 360

AAATGTATTA TAAAATAAGT TCATGCAAAA ACTTACAATT TTTCACAACA AACTACAATA 420

AAATCATGTA AACAAACAAT TTCTTTGAAA ATTAAGCAAA TTTATAAATA TAAATTATAA 480

AGATATATAT TTTTATATGA TCAATAATAA AAATTAATAG GATACTTATT TGGAAAAATT 540

ATTGAAAAAA CAATAAGCAT GAATTGCCAC AATAAGCTAA TTGTCACTTA ATAATTCTTG 600

TTTACTAGAC CACATTAGTA TAAACTCAAA TATTGGCTAC TATAATATAG GGGCTTTATA 660

CGCCACATGT TTAATGATAA CATAAGAAAA TATTGCAATA ATAAAAAGAT TGAAATATCT 720

TTATTAGAAA AGAATCTCGA TAATTTAGAA AACAGAATAA AAATCATAAC TAATAAATAT 780

AACGTTGAAA AAAATATATT CAAACTTTAA CTATACAATT AATTACACCT TAAAAATGCG 840

TTACATAAAA ATTAAGGACT ACTATAAATA GAAAACACCA CATAACCTAC AGACTCTAAA 900

GGAATAATTA AATCCTCATA TTTCAGTTCT CCAAAAGTTT AAATAGGGGC CTTTTACTTT 960

TCTTGATTAG CATATACATT ATTAAAGGCA TCTTCTTGGG CACTATCCTA AACTTTTTTA 1020

CATTATTATT ATTTTATTCT TTATTATTAC AAGATAATTC AAGAATCTAG ATTACAAGAT 1080

ATCAATCCTG CCATTAGTAG TTCAATAAAA CATTTAGAAT ATTTATACAT TATTTAATGT 1140

ATTTTTTTCA TTTTTGAAAT AATATTGTTA TAACTTAACT TAATAAGATA TTTGATTTCT 1200

TCAACTTGAG AATCCGATGT ACATAGAATC TGAACATCTC CTCTGCCCCA TTTGCCAATA 1260

TTCTTAATAT ATCTAGTAAA ACCCTCTTTT AAAATTATTT GATCTAGAGC AACAGTAATA 1320

GTAATATTAA TTTTATTTAC CCCAGGTCTA AAGCTAAAAT CTACAAAATA TCCGCCCTGT 1380

ACTTTAAATC CTGTATAGCA CTGTGTTTCA ACTTTCTCAA TTTCATTAAA ATTTAAAACA 1440

AAAATAAAAT CTTCTAATTC TTTATATATT GCTTTCATAT CGGAATTTAA TTTTTCAAAT 1500

TTTTTTAAAT TTTCGGTTTT AATATTATTA TCTTTTATAC CAGAATCTGT GTCATCTTCT 1560

ATGTCACTTT TCTTGCTGTT TACTAATACA TCGCTTTTTT TTTCATCAAA AAACATACTA 1620

AAAATATTTT TAATAATATC ATTAAATATT TTATCTGAAT ATGTTTTTTT AAAACCAATT 1680

TTAGCTTTAA AAAAATCAAG CAAATCAACA CTTGGATTTT TTGTTTCCTT TTTTAAATAA 1740

GCTGAAAATT TGTCTGTATA TTTTTTTTCT AATGCAAAAG ATCTAGCCTC TTCAACATTC 1800

AAAGAATTTC TAGAAAACTT TTTAAGATAT TCAAAATCCT TAGATGTTAA TTTTTCTAAA 1860

TTAACAACCA TAAAAGGCTC ATTGTCTAAC AAATTATCTT TATCTAGGTC AGTATAGAAT 1920

CTATATTCTA TGCCATCTGT TAATATACCA AATTCAACTC TCTTTGCTTG AGAACGAATA 1980

TTTTCAAAAT AAGGTTTTAA TTGCTTTAGA TGATTTTCAA GCTTTTCCCT GCTATTATGA 2040

TATTTGGCCT CTATTAAAAT AGTGGGTTCT TCATCCTTTT TTGTTGGATA AATAACATAA 2100

TCAACCCTTT TTAGTCCATC TTTAAGAATA TCTGCCTTCT CTTCAACTTT AACAATTGAA 2160

ATATCAGTAT GATCATAGCC CATCGCATCT AAAAATGGAT CAATAAGATT TTGTCTTGTT 2220

TGTGCTTCAT TTTCAATAAG ATCCTTATCC TTTTGAATTT TTCTACTTAC AGCTTTTATT 2280

GAATTTTCAA AATTTATATC TTTGTATTCA TTTGGCATAA TTATATTTTA CCAATAAAAT 2340

TAAAAATTAA TAATTCTAAA AATAAATTTC CAAAATGTTG TCTATTTTAA ACTCTTAACT 2400

GATACCTTAA TTCTTTTTTC TACCTAATTT TTTAGTTTAA AATCTTATTT TTTAATTTTA 2460

TTATTTTTTC CTTACCTTAT TTATACTAAA ATTTTTAGTA TTTAGCGAAT AATTTTCATA 2520

TCCTTTTATT AAAGACAAAA TATGATTTTC TCTTTTTTGT TTTTTAATAC CTTAAAATCA 2580

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ATTTTATTTT TCTTTACTAA ATTCAAAATG CTCTAAATAA AGCAAATTAG AGAAATTCAA 2700

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TTAATATTAT ATATAATAAT ATTATATTAT ACAAACAGTT TTTTAAACTA TTATTAATTT 910680

GATTTAATTT TTTTTATACT ATTAAATTAT ATTTA 910715

(2) INFORMATION FOR SEQ ID NO: 2:

(i) SEQUENCE CHARACTERISTICS:
(A) LENGTH: 116217 base pairs
(B) TYPE: nucleic acid
(C) STRANDEDNESS: double
(D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 2:

TAAATCCAAA CACTACAAGA TTCCATATAA CAAAAAAAAA TTTCAAAAAG AATGCTTTAA 60

AATTAAGATT TTCTGACTTC TTAAAATCAA AAATTCAAAA TGGAAAAATT ATAAAAGCTT 120

TCCAAATGAA AAATGAAAGA ATCATTTCTC TTGAGATTTT GCAAAAAGAT ATGATTATCT 180

TATTTATTAA ATTGTGGCCA TCCTCGCCCA ATATAATAGC TACAAATTCA AACTTTAAAA 240

TACTAGATGC ATATTACAGA AGGCCAAAAA TAAAAGAAAC AACAGGTGAA ATCTTTTTAA 300

AAGCTAAAGA AATACATGAA AGCAATAAAA TGTCTGATAA AAAAATTATG GAACTAAAGG 360

AAGAATATAA TAATACCAGC TATACATCTT ATTCTGAATT TCTTGAAAAT TACTACGAAT 420

CGCTTAATGA TCAAATTAAA AAAACCAATA TAAAAGAATT GCTTATTGAA AAATATAAAA 480

AAGAGTTAAT TGTTTTAGAA AAAAGAATAG ACTCTTTAAA ACAACAAATT AAACTGCTTG 540

AAAACATTGA AAACGAAAAG GAAAAAGGCG AATTGATTTT ATTAAATATT AACAAAATAC 600

AAAAAGGGAT TAAAGAAATA AACCTCTTAA ATTATAAAGA AGAAAAAATT AAAATATCCT 660

TAAACCAATC ATTATCACCA AAAGAAAATG CCTTGCAATA TTTTAAAGCA TATAAAAAGG 720

GCAAAAATTC TTTCAAAACC ATACAAAATC AATTAAAAGA TAATCTAGAT AAATTTAATT 780

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CCTATTGTGG ATTTGAAATT CTTATTGGAA GAAACGCAAA AGAAAACGAT AAACTTTTAA 960

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AAATGTGTAT ACTAAAAAGA AAATTAACAA TATGTATGAT ATTTTTATTT TTAATTT 116217

(2) INFORMATION FOR SEQ ID NO: 3:

(i) SEQUENCE CHARACTERISTICS:
(A) LENGTH: 111309 base pairs
(B) TYPE: nucleic acid
(C) STRANDEDNESS: double
(D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 3:

GTATCTATTA GTAGATTTTT ATATCTATTT GTTAGTGGGT ATAAATTGAG TTCGTTTTTA 60

AACTCGTTTA AGTATTTTAA GGCAACAATA GTTGAATTTT TTATTGATTT TGATGAGTTT 120

ATCATTTCTG TTGAGTTTAA ATATTTCTTT TCTTGCTTTG GTAGTTTTGG TATTTTTTAT 180

TTGATTGAAT TTTGAAATAA TTTTTGGTTC AAACTTTTTT TCTTGTAAAA AATAAATTAT 240

TGGCAAACTT TTTTTCCCTT CGAGTAAATC ATCTCCGAAT TCTTTACCAT TAATTTTATT 300

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GTAAATTTTT TTAGCTTTGT CTTCATTATT TGTGAGTATT GCAGCTAGAA AGCTGGCCAT 420

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TAATTTTGCA GGTAAAAAAT AAATTAAATT GCCAGCATTT ATACTGTTAT CTATTCCATA 660

GATTAAATGT ATTGCTGATG CACCGCGTCT TTTTAGTGAA TTGTCTTCAA TGTCATCAAT 720

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TGATTTTCTG GTCGATATTT TAAATTTTTA TTTAAAATAT TTAATATTTT AATTAAnTCC 111300

GGnTCCTAC 111309

(2) INFORMATION FOR SEQ ID NO: 4:

(i) SEQUENCE CHARACTERISTICS:
(A) LENGTH: 53585 base pairs
(B) TYPE: nucleic acid
(C) STRANDEDNESS: double
(D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 4:

AGCATTTAAT TTACAATTTA AACTATCTTT TAAAGATTAC ACCTTATGCT AAAAAATAAG 60

TTTTTGGTCG CAAAAATCGC AAATCCACAA TTTATCAACT AAATACTAAT ACACACTTAC 120

AATCAATCAA TTCCTAGCTT ATCTTTCAAT AAGTTCAATT TTTCAGCATC CATACTAAGC 180

TTATTTAAAT CCAAAGCATT AATAATTGCA ATTATCTCTT CGTGGGAAAA TCTCTTTCCT 240

CTTAAAAAAT GATTTTCAAA ACTGTTAAAA GCGATGGGCA CAATTTCTCT TACTATGCTT 300

ATCAATGCTT TAGCATATTC GCGCATTTCT TTTGGCGAAT TTTCTTTAAT TTCTTTTGGA 360

CTGTCTAGGG CTAATCGCAA TTTAATAAAA TGAAAAAGAT TATTTAAATC AATTTGCCAA 420

TACCATTCGG TATATAAACT TAAGGGCAAA ACTATTCTTG AGAGTTCTTT TGGAATATTA 480

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GACAATTTTT CCAAAGACTT AAACTCTTTT TCAGAGCTAT TACTAGAAGT TTGACACTTT 600

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TCATTAATCC TTGCCGTTCT ATGCCTCATC CATTGCCTTG CAACAAATAT TGGAGCTTTA 720

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ATTAAAAATA AAAATTGTAT TATGATTGGC ATGGGCTTCC tATTTTATAT TAATATTCTT 53580

TAATT 53585

(2) INFORMATION FOR SEQ ID NO: 5:

(i) SEQUENCE CHARACTERISTICS:
(A) LENGTH: 35515 base pairs
(B) TYPE: nucleic acid
(C) STRANDEDNESS: double
(D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 5:

GGTTTAGGAT CCTTAAATAC GTTGATCATG CGAAATGACA TATCAGAAAG TCTTTTCCTA 60

ACACCATTCA TTAGACAACC CCCACACAAG ATGGCGTTGA AGTTTCTCTT TTTAGTTTTT 120

CTAAAAATGC ATCAAGTTGT GAACAAAAAT TCTTGTTTGA GCCACAACTC CCCTCGCCGC 180

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TATGGTAGAA AGTAAGTAAA AATATTTGTT CATACGTCAG TGAACTGGCA TCAATACCGC 360

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AGAAGATAAA ATGCCACTTC CCCTATCATT ATTrCTCTGA GTAAGTGCAC TTAAAGAAGT 35460

TGTTGCGCTA GACAGTGCGT CTTGTAGTTG TACTAAAGAT TCATCTTTGT AAAAC 35515

(2) INFORMATION FOR SEQ ID NO: 6:

(i) SEQUENCE CHARACTERISTICS:
(A) LENGTH: 26811 base pairs
(B) TYPE: nucleic acid
(C) STRANDEDNESS: double
(D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 6:

GTTGTTTATA CATTATTACC TTCTTTGTAT AAAATCTATC CAATAGAAAA GTTTTAAAAT 60

TTTTCTATTG GATTAGCTGT TTTTATATTA TATTTTAAAA ATTTTACATA AGCTTGTCGA 120

TTTCCATTTT TATATTATCG GACAATCCTC CAAATACAAT TTGAACACTA TTTCCTTTTT 180

TTAATATTCC ACTGGCACCT AGATTTTTGA AATAGGCGTC TGATTTAATG ACTTCTATTT 240

GTTTTAGATT GACTCTTAAT CTTGATGCAC ATGCATCAAT GTATGTAATA TTATCTTTTC 300

CTCCAAGACC CTCTAATACT TTTGAAGCGG TTTCTGATAA GCTTGTTTTT TCTGAACTCA 360

TCATGGTTTT TTCCATCTCT TCGTCTTCTC TTCCTATTGT TTTAAGATTG AATTTCATGA. 420

TTGCAAATTT AAATATAAAG TAAAATCCAA TAAAGTAGAA GATTCCCAAG ATAGGAATTG 480

CTATCCAATT TGTTTTGCTA TTTCCCTGAA GTATTCCAAA TAGGAACATA TCTATAAATC 540

CGCCAGAAAA TGTTAGTCCA ACTCCTACGT TTAAAAGGTG TGTTAAAAGA TATGCTAGTC 600

CAAAAAGAGG AACATATATA AAGTAATAAA GTGCTGGTGC TGCAAAAAGG AATGCAAATT 660

CAAGAGGCTC TGTAATTCCT GTTAACATTG ATGTAAAGCT AGCAGATAGT AGCAATGAAG 720

CTACGTTTTT TTTATTTTCA GGTTTTGATG TATGGTACAT AGCAAGCGCC GCCCCAGGTA 780

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AGTGTACAGT ATTGGGATCT GCAAGTTGTT TGAAAAATAT ATTTTGTGCT CCTGAGATTA 900

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GATGCACATG CATCAATGTA TGTAATATTA TCTTTTCCTC CAAGACCCTC T 26811

(2) INFORMATION FOR SEQ ID NO: 7:

(i) SEQUENCE CHARACTERISTICS:
(A) LENGTH: 26782 base pairs
(B) TYPE: nucleic acid
(C) STRANDEDNESS: double
(D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 7:

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AATCTTAGAG TCGTATTTTT AATGTGACAA CTACTGCATA GTTTACTTGA TGGAAAATCT 120

CTATCTACTT TATACAAAGA GGATCCATAC CACTCTGATT TGTATGATAA TTGTCTTACA 180

AACTCAGACC ATCCTAAATC ATTAATACTT TTTCCAAACA TTCCTTTTTG CATGCCTTTA 240

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TGCAAAAAGT CTTTTCTTTG ATTTGAAATT TTCTTATGCA GCTTAGCAAC TCTTAATCTA 360

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TTTTGGTTTA AATGTTAACC CG 26782

(2) INFORMATION FOR SEQ ID NO: 8:

(i) SEQUENCE CHARACTERISTICS:
(A) LENGTH: 18359 base pairs
(B) TYPE: nucleic acid
(C) STRANDEDNESS: double
(D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 8:

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GCGACAATTA GCGATCCTCA TGAAATAGTA AATGTTTTTT TCCGCATTTA AAAAAGGCAG 13860

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TGTTGGGATA TATTCTGCAC TTATAAGCTT TATTAGCATT CATTATTTTT TATAATTAT 18359

(2) INFORMATION FOR SEQ ID NO: 9:

(i) SEQUENCE CHARACTERISTICS:
(A) LENGTH: 14752 base pairs
(B) TYPE: nucleic acid
(C) STRANDEDNESS: double
(D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 9:

GCAGGTCGAC TCTAGAGGAT CCCCTTACTT ATAATAGCTA TATACTACGA AATATACAAA 60

AGAAAAATAT ATAAATGCCG TTTATATAAA ATTACTTGGA TTAGAAAAAC AGGGAAAATT 120

ATTTCCTATA AACCATCAAA AAATATAAAA ACTTTCAAAA AGTATGAGGG AAAAAGTATT 180

AATTTACATC CCCCCAAAAG AAAAAATTTG AAAATATTTT ATAACACCAA ACTTTTAAAA 240

AGTAACAACC CTAACAAATA TTTTCTCAAG TGCAATCTAT TAAAATTGGT CAAAATGCCA 300

TAACGTTTGA AGAGCTTGCA ATCAAATATA TAAAAATCCA ATAATATTGC CTAGCAAATC 360

TTATTATTTT AAAATTTTAA AAAAACAAAT CCTAACTATG TCTTAAATCT GGATCAATAT 420

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AACAAAAAAA TTTTTTGTGT GGAAATTTTT CAAAAAAATA ACATAGTTAA AAAGGCATTT 600

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GATAAACAAA AATGGAATTT TTTAAAAATT GAGATAGACT AAAATTGAAG GAGAATTAAA 720

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CTTTGTTTTC ACTATAGAAC AATTCTTGAT TCATGTGATC TAGTGTTTTG AC 14752

(2) INFORMATION FOR SEQ ID NO: 10:

(i) SEQUENCE CHARACTERISTICS:
(A) LENGTH: 10749 base pairs
(B) TYPE: nucleic acid
(C) STRANDEDNESS: double
(D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 10:

CTTTTTTACT TTTAAATTTA GTAAATTGTA AATTTGATAG TCTTAATTTA TCTACAAAAA 60

GCGTAGATGA CAAAAACAAT TCTATAGCCA AGCTTCTTCA ACACTTATCA AAAAGTGAAG 120

ACCAAGCCAA TAAAACTTCT ACCTCAGAAG ACCAAAAGGA ATTAGAAATT ACGGAAAACA 180

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ATATTTACAA TTCTTATATC CCAGAAGTAA AAGAGGAAAT TGTTTATGAA ATTCTTGAAG 360

AAGTGATAAT TCCCGAAACA AAAATTCCTG AAATTACTGA AGAAGTGATA ATGCCTATTC 420

CACAAACAAT AGATTTTTAT ATTGAACCAA GGCCAATAAG TAGTTTCCTT ACTCAAGGGA 480

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TTGATCCTGC AATGCCTTGT AAGGAACTCT TTGGCCTTAA ATCAGAATCT ATAGTAAGAT 9420

CAATTTCATT ACTTTCAATA GGCTCCTCCA ATTCACCACT TTCACCGCTA AGGGAATTTA 9480

AACTTCCTTG AGTCATCCCT TTATATTCAT CTTGTTGTTT TTGTATTTTT TCTTTTCTAG 9540

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TTTGTAAAAA ATTGCTTTTA TTAGCAGAGT CCCTTTTTAA ACTATGAATA TACGGTTCTT 9660

TTCTAGAAAC CTTGCTTTTG CCTTTCTTTC TAGCAATCTT ACCTTTACCT TTCTTATTCA 9720

TAGGCTTTTT AATAGATTCC TCGCTAGTCT CTAAAATAGA GTTTCCCTTA TCAAATAAGC 9780

CAGGGGATTC CTCTTTCATT TCATATCTTA TGAATAAATC ACAACTTATA AAACCAAAAA 9840

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AATACTAACT TTAAATCATT TTAAGTCGGA TTCCAAAAAT TAATAAACTT TTCTTATCAA 9960

AAAGAGGCAA AATTATTACA ATTTTAATTG TAATAATACT AAACTAATAT TTACTATTAT 10020

TTATCAATTC ATTGAATTTC TCTTTCTTTT TAAAAATTTA AAATAAGTGC TATCAAGAGG 10080

GTGAACAAAG TACTTTTTAT TAAATTTTAT AAACAGATCT TAAAAAATAA CTTCTAAATA 10140

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TTCGAACTGT AATCAGACCT TTATATTCTT TATCAATTAT CATTGCCCTG AATCATAAAT 10260

TTTGCTTTAA AACTTGATGT GCAGGTTTTA GTATTACTAA GTTGAAATAT TGCAAATTTA 10320

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AGAAAATTAG AAAATTAAGT AGCTACTATA AAATTTTTCC ACTATGGAAT TGAGTTTCCA 10560

GATGTTCAGG AAGGATTGAT ATTGTAATTA GAAATCCACC AGGGAGAAAA CTAGTTTAAT 10620

GAATCAGAAT CTTCTCAAAA CATATTCCTA GCTACAGAAG ACTAAGCATA nAAGACCAAG 10680

TAAAATAAGG CAGAAATACT AGTAAGACAT TATATTGAAT ACTGATACAA TGAGGAnAAA 10740

GGCGTATAG 10749

(2) INFORMATION FOR SEQ ID NO: 11:

(i) SEQUENCE CHARACTERISTICS:
(A) LENGTH: 10502 base pairs
(B) TYPE: nucleic acid

(C) STRANDEDNESS: double
(D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 11:

AATGAGCAAA GTTTGATGGA ACAAGAAAAG AACTCTCTAT ATGTATATGC GCATCAATAA 60

ATCCTGGCAG CACATATTCA TCCAATGTCG CATTAATTTT TTCTATGCTC GCAATATGAC 120

CGTTTGCAAT CGCTATACTA GCTGGATAAA TTTCTTTATT AAAGATGTCA ATATAATTAG 180

CTTCAATTTT AAATAAATCC ATTATTTATC CTCTCCTAAA AACTTTTTGT TCATAATATA 240

CTCTACCTTA AAATCTTTAA ACTTAAATTA TAGCGCAAAG TGCCAGTATA TTTAATTGCA 300

ACTGTATTTA GAAAAAATAA AAATCGTCGA TATTTGTTAT ATACGATAAA AATCGAAGAT 360

AATTTTTTGG GTAAAATAAA TAGAAAAAAT TTAAATAAAA AATAGCTAGC AATTCTAATT 420

TTGATACCAG CAAAGATTAT GTGGTAAAGT ATCAAACAAT TTTTAATTGA TTTTTATAAA 480

AAGGTTTTAC ATTAAATTTA AAATTAAATA ACAATAATAT TTCGAATCTC TTTTTCTAGA 540

TTTATAGGAT TTTTATCCAT GCTTAAAACA ATATCTGAAA AAGCTTTATT TAAATAATCT 600

CCCAATTTGA TATTTTCAGA CTTGATGGAC TCTTTATTGC TGTTATAATC ATTTAATAAT 660

TGTTTAACAA GCTCTCTAAA GGTTGTTCTT AAAGATAAAA AATTTTCAAG ACAAATTTTA 720

ATCCTTTTAA GATCAGGAAT ATTTAATAAC ATTAGGTTGT TTTTTTTATC AAGTAACAGC 780

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GCAATCTTCT TAATGCTATG TAAATTTGAA TTTAAAATTA ATCCTAATTC TTTCAACTCG 900

CTTTCATTGT AATCTAAAAC CGCATACAAA CACCCTCTAG CCTTTTTAGA TTCTTCTACA 960

TTATCAGCAA GTGTTTTCCT TAGTGTATAC CAACTAACAT TTTTAAATGG AAGATTATAT 1020

TGATTAGATG GCTCACTTTT GAATTTTTCA AAAGAATTAT CTATGGCTTT ATTATTAACT 1080

TCTACAAAAT TATTTATCCA AGATAAAAGT TCATTCTTTT CTCCCCGCAC ATATTTTTTC 1140

CCAAGTTGTT TTTTATAATT ATGATTTTTG CTAGCCTTAT TTTTATTTAA ACCATCGTTT 1200

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CCCTTAAAAT GGTTTTACAG AATATCTGCA AAGCCATAAT ACAAACTTAT AATAAATATT 1320

AAATAGTAAT CATTATAAAA TTAAATATAA TTTAAAAATG ATTTTTTTGC AATATTTTGA 1380

AATGTTAAAT AATTATGTGC TATTTCTAAA CACAAGATTA TAAAATAACT TCTTGCTTTA 1440

ATATCATGAT GGCTTTTAAT TGCTTCAATT AACTTAAATT ATGCAGCTAT CTTGGCTTTT 1500

CTTTTTTATA AAACATTGTC TTTAGCAAGG TCATAAAATT ATGATCTACT AAAATGTATT 1560

CGGATTTATT AATATTAAGC AATATGTTAA AATAAATTTA AATACAGTAA TTTAATATAT 1620

TATTGCTTTT AGGAGGTCAA TCTATGTTTA GAGGAAAAGA AATATATTTC TTGTTATTTT 1680

CGTTACTTTT GTTTATATCA TCAATTATAA TTTCTCACGG AATAAAAAAC ATTGGCACCA 1740

AAAATGAAAA TTATATTACG GTAAAGGGTC TTAGCGAAAG AGAAATTTTA TCAACGTCTT 1800

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ATAATTTAAG CTTATCGAAG ATTAAAAGTT TTTTTTTAAA ACATGGATTT AGCGAAGACC 1920

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AAAACGCAAA TCAAGGATAT TTTGAATTTC TTCCAATTGA TAGAAGCTTG GGCGATCAAG 2280

AACGTTATCC AAAAAAAATA TTAAGAATCG TTACAACCGT TTCTTATTAT TTGGATTGAT 2340

ATTTGGCGAA CTATTTTTTC TAAAAACCCA TATAAATAGC TTGCGGTTTC ATACCATGAC 2400

GCGATCTACA CACCTAATTG AAGCATAGCT ATATATATTT ATTGCTGTAC TCTTTATCTT 2460

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AAGTTCGCTG CATACACCTT CAAAGTATGT TTCTACATTT TTCATAAGAG TTTGAATACA 2580

TTTTTTAGTT TCAATTTTAC TGTTTGAATG TTCTTTAAAA GAAACGCTTA CCGAATTTTC 2640

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GATTTTTTCC GAACTTAGGC CTGTGGCTTC ACTTATCTCA TCTAATACTT GATCAATAGA 3060

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CAAAGGGTTC TTGCAACTAT TTACTAAATG TGCGCCCTAT TTGTGAATTT AAAAAAATCT 3840

CATAAATGCC TATTAATTAA AAATATCCAA AAAATTGAGG TAAAATGGGG CAAATATTGA 3900

AAAGTATATT CCCAATATCC CAATCATAGT GCCAAAATCG ATTACTAAAA ATAAACCCAT 3960

GAACTGCTTT ATAACAAAAA AAATATTTTT TCCAACATTG CTTTCATAGA TATATGGACT 4020

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CGAACTCAAA GATAAAAAAG ACGACAAATA TACTAAAATA TTGGCAGCTT TGAGTGAGGC 5160

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GGTATACCAA GGATGAATTT TTGAATATTT TGGATAACGA ATTTAGAGGT AACTGCTTGT 7380

CAGTTGTAAA AAAGATTAAC GAAATTATGA CTATTTTAGA AGATATCCAT CAAAAAATCA 7440

AAGGTATTAA TTATACCATT AAAGAAGATT TAGAAAAATC TAAAAAGATA TACAAAGAAT 7500

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CAAGAAGGAT TTGATATTGT AATTGGAAAT CCTCCATGGG AGAAAACTAA GTTTGATGAA 7800

TCTGAATTTT TATCAAAACA TGTTCTTAAC TACAGAAAAC TAAGCATAAA AGAACAAAAT 7860

AAGATAAAAC AAGAACTACT TAGTAAAGAG AATCATCCTT TGAGTATCGA GCGCTATGAG 7920

GAAAAGAACA GTTTGAAAAC TTTGAATAAT ATGTATAGGT TGGTGTTCAG AGAATTTTCT 7980

AGTGGGGGGG TCCGAATCTT TTTAGATATT TTACGGCTTT CAATTTGAAG CTAGTGAAGC 8040

CCGGCGGCAA CTTAACTTAT TTGACTCCTT CCAGTTTGTG GAGCGAATCC GGCTCTAAAG 8100

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TGAAAGAAAT AACCAGTGAT CTGAAAGAGG GTAATGAAAA TGCTTATAAG GGAATCGAAT 8340

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TGATGAGCCT CTGTTTGTGA AAGTCTGAGT TTGGTACACA CAGACACTGA AATTATTTAA 9360

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GTTATTTTTT ATTGAGACTT TCTAAAACAG TTTTTGCATG TTTAATAAGC TCTTTTATTT 9480

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TAGATATTTA AGATTTTTGT ATAACAAAAT GTTAAGTGAA AAGATTATAT AAATTAGTAG 10500

TT 10502

(2) INFORMATION FOR SEQ ID NO: 12:

(i) SEQUENCE CHARACTERISTICS:
(A) LENGTH: 9842 base pairs
(B) TYPE: nucleic acid
(C) STRANDEDNESS: double
(D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 12:

GAATTTAGTG ATATTGCAGA TATAATCAGT TTAATTGAAA ATATTTTGGG GAATAAGCCT 60

TCTAGTGGGG ATAAAGAAAA AATGAAGAAA ATTAGAAAAT TAAGTAGCTA CTATAAAATT 120

TTTCCACTAT GGAATTGAGT TTCCAGATGT TCAGGAAGGA TTTGATATTG TAATTAGAAA 180

TCCACCATGG GAGAAAACTA AGTTTAATGA ATCAGAATTC TTCTCAAAAC ATATTCCTAG 240

CTACAGAAGA CTAAGCATAA AAGAACAAAA TAAAATAAAG CAAGAAATAC TTAGTAAAGA 300

CAATTATTAT TTGAATACTG AATACAATGA AGAAAAAAGC AGTATAAGCG CTATTAATAA 360

ATATTTATAA AAGTGATTTT AAAGACTTTG CTAGTGGTGG GGATCCAAAT CTTTTTAGAT 420

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CAAATAATCC TCCATTAAAT ACGGGGAACT TTATTGAATC ACTTCCCTTA TCAAGTAAAT 960

TAAAAATTGT TATTATTTTT TATATCCTAA TTTCTTTTTT GTATTTTAAT CATAAAAAAA 1020

TATCTAAAAG ATATAGAAGA TCTGTATAGC TTGTTTTCTT CTAAAATTTT CTTAAAACTG 1080

TCGTTGTCTT AAATATATGC AATAAAAAAT ATTCTTAAAA TAAAAATAAT TGCCTCTTTA 1140

AGTATGCTAG CCAAAATATA ACGAGTAAAT TCTTTGTCTG ATACTTTAAA TTCTTTGTCA 1200

TATATATTTT TTTTACAATT TTAAATACTA TAGAGTCATC AGGCTTCTCA TAAAGCATCT 1260

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CACATATTCA TCCAATGTCG CATTAATTTT TTCTATGCTC GCAATATGAC CGTTTGCAAT 9480

CGCTATACTA GCTGGATAAA TTTCTTTATT AAAGATGTCA ATATAATTAG CTTCAATTTT 9540

AAATAAATCC ATTATTTATC CTCTCCTAAA AACTTTTTGT TCATAATATA CTCTACCTTA 9600

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GAAAAAATAA AAATCGTCGA TATTTGTTAT ATACGATAAA AATCGAAGAT AATTTTTTGG 9720

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CAAAGATTAT GTGGTAAAGT ATCAAAGGGG ATCCTCTAGA GTCGACCTGC AGGCATGCAA 9840

GC 9842

(2) INFORMATION FOR SEQ ID NO: 13:

(i) SEQUENCE CHARACTERISTICS:
(A) LENGTH: 9542 base pairs
(B) TYPE: nucleic acid
(C) STRANDEDNESS: double
(D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 13:

TGAAAACCCT AAGGATGAAC TTGCCGGGGA TTGATAAAAA TCTTAAAGGT TATGGCTATA 60

AGTATCAGAA TTTCAACGAA ATAGCTAGAG AAATTAAAAA AGTTATTGAT AAGCACAATT 120

TATGCCTTGA TTTTAAGCAA TTTCCGACTT TTACAGTTGT GGGGGAGCAA CAAGTTCTAC 180

ATGTTGTTAG AACTACGTTT TATAGTACAA ACACTGGGTA TAAAGACTCC TTTGATACGC 240

CAATACTTAC AGAAAATTTA AAATGGAATA ATGAAAATGG GTCTAAAAAT GTTGTAAATA 300

CAGTGCCACA ACTGGTAGGC TCATCAATTA CTTATTTTAA AAGATACGCA TTAGTAGCAT 360

ATCTTAACAT AGAAAGTGAA GTGGATACTG ATGCAGCTCC TATTTACAAT AACCACGAAA 420

ATGAAAATTC TATGCCTAGC AAACAAGCTG GTGTTAACCA AAATCAAGTA AAAAATTTTG 480

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TTTACTAAAT CTTGCATGAA AGAAAAACAA GAAAAATTAA AGAAAATTCT ATGCAACACT 7380

CAAAAAGAAT TTGAAAAAAA TGAATATAAT CCTAAACAAT TAGAAATAAG TTTCCAAAAA 7440

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TAAAAAAGTT ATTGATAAGC ACAATTTATG CCTTGATTTT AAGCAATTTC CGACTTTTAC 9540

AG 9542

(2) INFORMATION FOR SEQ ID NO: 14:

(i) SEQUENCE CHARACTERISTICS:
(A) LENGTH: 9399 base pairs
(B) TYPE: nucleic acid
(C) STRANDEDNESS: double
(D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 14:

TTATTACATT CTTTTAATAA AGATTTACGA TTCTTAATGC TTGGATCTAA GCCTTTTTTA 60

AAATCAATAT ATCCTTCACC AAGAGCACTA AATTTGCTAA ACATTTTATT AATAAGAGTA 120

AATTCTTCAT TGTCTTTGAA TTCTATTATT GATTCTTGAA TAGGAGATAG CTTCTTAATT 180

TGATTTATAT TTAATTCAAT TCCTTTATAA GCATCGTCTT TACTATCTTT CAAATCCCTA 240

GTTATTTCTT TTAAAATATT ATCACTACTC TGAATCATGA ATTTTGCTTT AAAGCTCGAT 300

GTAGATTCTT TAATATTACT AAGTTGAAAT ATTGCAAATT TAAAACTTGA ATGCACATCT 360

TTAAATCTTT TTTTGTTTTC AAATTGATAA ATATAGTTAA GTTTATAGCG AGCAAATATA 420

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TCTTGTCTTA TGATATTTTG TTCTTTTATG CCTAGTTTTC TGTAGTTAGG AATATGTTTT 720

GAAAAAAACT CTGTTTCATT AAATTTAGTT TTCTCCCATG GAGGATTTCC AATTACAATA 780

TCAAATCCTT CTTGAATATC TGGAAATTCA ATTCCATAGT GAAAAAATTT ATAGTAGCTA 840

CTTAATTTTC TAATTTTTTC TATTTTTTCT TTATCTTCAC TAGAAGTTTT ATTGCCCAAA 900

ATATTTTCAA TTAAACTAAT TACAGACGCA ATATCACTAA ATTCCATATT TAAAGATTTA 960

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ACTAGTATTT CTTCAATCTC GCTAATACTG AGCAAACCTT CATTATTTAA ATATTTAACC 1980

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ATTGATTCTT CAAGAATGCT AGCTAAAATA TGCTGAGTAA TTTCTTTGCC TGATAATTTA 2280

AATTCTTTGT CATATATATT TTTTGCAATT TTAAATACTA TAGAGTCGTC GGGTCTCTCA 2340

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ATGTTATCTA TATTTGTCAA CTCTTTATTC ATATGACTGA AAGTCCTACT ATTTAAGACT 9360

AGCTCTTTTT TGTTGTATTT TTTCCCCATC TTATCATCC 9399

(2) INFORMATION FOR SEQ ID NO: 15:

(i) SEQUENCE CHARACTERISTICS:
(A) LENGTH: 9360 base pairs
(B) TYPE: nucleic acid
(C) STRANDEDNESS: double
(D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 15:

GGGGTACAAC AGAACGTAGA AGTTGAGGCT GGTGGTGGGG ATGACTTGTA CGGAATATGC 60

ACTGATATAG ATGAGTTTAC TGGCATGGCA ACTGTAGTTC CAATTACAAA TAACTTCACA 120

GGGTATTTAA CATTTAAGAA AAATGGACAA AATGGTGTGA ATCCGGGTGA TAAGCTGCAT 180

TTTAATGCAC AAGGAGAGCT TGAAAAGAAT GGGGGAAATG ATAAATCTGT TAATGCTATA 240

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CTTTTTATTA TCCCCGTGAA AACCCTTTCA GCAGAAATAA TACAAGAATT TTTAAGCCAG 9360

(2) INFORMATION FOR SEQ ID NO: 16:

(i) SEQUENCE CHARACTERISTICS:
(A) LENGTH: 8905 base pairs
(B) TYPE: nucleic acid
(C) STRANDEDNESS: double
(D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 16:

AGTCTGAnGT GGGCCCTTnA ATATAAATAA AAAGGTTGAT AAAAAAATCG AATAGCGTCC 60

ATTAAAATAA ATAATTCTAA AAGTTCTTTA GATTTTACTA ATATTTTTTT AAACTTTATG 120

GGATATAAAG TTAAATTTTA TTTATTATGC TTATGATCTA TATTCTTATC ATTTGGTATG 180

AAAAAATTGT AATTAAAGAG GATATTAAAT ATTAAATATT AAAATATATT AATAATAAAA 240

TTAAGTAAAT TTTTATATCA ATGGTAGTAT TTTACAATTT ATTTTTGAAT TTCATTCAAT 300

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TCCGATAAAT ATTTGAGTAA AGAATCTCGA AAATATAACC GAAGCAAGGA GCCATTTCTT 7620

TAATTTTTGA TTTCTAAATG GTATTAATGT TTTTTATTTT CATTGATAAG CTTAAGAAGA 7680

GGGTATTAAT ACTAACATGC TCTTCCTAGG TAAATTGATT TTTAATAACT TTTTTTAAAC 7740

GACAGCTGTA ATCCGTTTAT TCTAATGTCA AAATTGGATT TAAAACCTAC AAGCCCAATG 7800

CCGAGTCTTT TTTTAAGTTT TCATTGTATC TGTTTGCAAA TTTAAAGGGA ATAATAATTC 7860

TAGTTGTGTA GGATGCTGTA ATCGTGAGTT CTCCTTCCTA TTCCTAATGA TTCTATTCCG 7920

ATTGCTTCTG TTGGTGTTTT TGACGGGATG CTTCTGATAA TGTATCCAAG TTATCTCAAG 7980

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TACGGGTGCT TTTTCGCTTT CATATTTCAT TACGGCTGCA ATATACTCAA CACCTTTTTC 8160

AACTCTAACT TATTATCTTG TGTTGCAAAG ATTTGCATTG TTAAACTAAA AATTAATAAT 8220

AATATGAAAA TTTTTTCATA TTATTATTAC TCCCCTAATA ATAAAGTTTT GATAATTAAA 8280

TATTAGTGCA AGTTATTAGA TTTTGAGCTT GTAGATTAGT TTTTCTCTAA AATATTATTT 8340

GAATTGCTTT GCTGTGCTAA GCAGATTTTA ATTTAGGTTT TATTAATTTT CCAGTGAACT 8400

ACTATTTCTA TAATCTTTGA TTATAGAAAT AGTAGTTCAC TATTAAGCCT ATCCTTTTTA 8460

TAAAGGATAG GCTTTTACAT GATATCAATT TCAATAACAT AACTTTAAAG GTTGCTTATT 8520

GATAATATGT TATTTTTCAG CTCTGTTGTT TGGAAGCTTT TTTTTACAAT TTGCTTGTAA 8580

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TTTTTAAGAC ATAATAAGTG ATTAATCACT GTGTCAAAAG TGCTTCCGAA GTTGCAAAAG 8820

GTGCCAAATA GGAGGGTTTT TTGTTTTGAT CGAATTAAAA TCTTCGAAAG ATTCGCTAAT 8880

TTATTGGTAT TAATAGAATT TAAAG 8905

(2) INFORMATION FOR SEQ ID NO: 17:

(i) SEQUENCE CHARACTERISTICS:
(A) LENGTH: 8318 base pairs
(B) TYPE: nucleic acid
(C) STRANDEDNESS: double
(D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 17:

CTCTACCTAG GACTTTAATT AAAAAATTTG GATTGTnATT CCCAATGCTT TTTCTATCTT 60

GnAAAGAACT TTAAAAGTGC TTTCAAGATA GTTTCTTTTT GCACGTTACT TGAATTTACA 120

TTCAAATAAT AGGGCAAGTT GCTTTAyATA TACTCTTTAT TTTTTTTGAT TTTTGTATTT 180

AAGTGTTGAT ATTATTAGTA ATATTTTAAA CTTTACTCTT AACTAAAAGC TTGTTTTATT 240

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CTATAAAAAT ATGTAAATAA TAATTTATAA ATTAATAAAA AACATATAAG GGAGCTTTCT 360

TAATGAAAAT CAAAAATATA GCAAACATAT TAATATTTTT TTTATCTATA GTTTTAAATA 420

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CTCAAGTAGC ATACGGCTTT AAAATTCCCT TTATAATCAT CCGGGGAATA TCTGATATAG 1080

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CAGCAAAAAT AGTAGAAAAC TTAATTAAGT TAATGTAACT TAATTAAGTT TTTAATTAGA 1200

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ACACTTATGC ACAATGTTGA AACATATTTT GAAAGCGTAT GTGATGAGCT TAAAGATAAA 2040

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AATTCAGCAA TAGCCATACA TGTTTGTTTT AATATAGATT AATTTTAAAT GCTTTTTATA 2160

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TTAAGTTATT GCATAAAGGA TTAATTTACT ATGTCTTGTC AGCAGTTCTT TTATAAGCAA 2280

CATTCCAGCA AGATATCAAT ATAAAAACAC CCAATATGAA AATATGCTAT TTAGAAAGAG 2340

GAAGAAGAAA AAGTAAAAAA GATAAAATAT TTTAAACCAC TTTCCATTAT GCTAAACTTC 2400

AATTAATCCA CATGGCTTAA TGATTCACTT TAAATGGCGG CTTTGAATTT CAAAAAAGCA 2460

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CTACGAGTCA CAAAATTCAA AAAATAAAAG AATCAAACTG CTATAATAAT TTTAAACAGT 2940

GAGGAAGAGT TAATATGATT AAATTCCCTA AAAACCATAT ATCAAAAATA CATATCATAA 3000

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AACAAAATTA ATTATGTATT TATAATAAGT TGTTTTTGTA AAAmACAAAA CCTCAAAATT 7080

CAAAAACCAC TATAATTTAA AATATGCATT CGAGAACTCA AATGTAATCA AGTTTACTAA 7140

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TAATCATTAT TAATAGATAT TAAATATCAT GTCCAATATT ATATTAAGAC ATGCAACTAA 7560

CATCTACATT TTTATATAAG CAACTCTACG GCCAAATAGT TAATACAATA TATCAATAAT 7620

GTTGTCAAAT TAAATTCCCA TTCAATTAAA ATTCAAATAT TTATACCTAT ATCCTTTAAA 7680

ATTTTTGACA ACGCTATTTA ATTTTTTTAT AAACTTATAT AAGCTTTTAA AGCTTTAATT 7740

CCCTCTTATT GAATTTTCCA GATCTAAATT CAGATTATTA TTAACAACCA AATTGCTCTA 7800

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CAGGCTAAAT TGCAATCTTC CCAATTATAA ACATTGCAAT AAATTATTAA TTTTTATGAT 8100

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CAAAATAACT AGGAGTATTA ATATGCAAAA TGCATTTCCA ACCGCAAAAA CACACCTTGc 8220

CAGAATAAAT TACAACACAA ATTAATAGTT TTTATCTCTA CGCTAAAGTA TATAAACAGT 8280

AAATATAAAA GATATACCCA AAGTAATATA CTTTATTA 8318

(2) INFORMATION FOR SEQ ID NO: 18:

(i) SEQUENCE CHARACTERISTICS:
(A) LENGTH: 7624 base pairs
(B) TYPE: nucleic acid
(C) STRANDEDNESS: double
(D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 18:

nTTCGCTTTA AATTTAGTTT AAAAnAATAA TAGCTATATT TTCAAATTAT AAATAGTAAT 60

CTATTTTATA GCGCTCGACT AGCTGGCAGG ATGACTAAAA AATTAATATT TTTTAAAAAA 120

GAGTGTACTC TCATTTATTT TGCCATATCC TCTTTAACAG ATATTTATCA AGAAACCAAA 180

AATACTTGCC TTATAGTCTC GCGCAGGAAA TTTCAGGACA TTTAGATATT TTAACCATCA 240

TGATTTATCC TTATATATTA CCCAATATTA AAAAGAAGTT CTCATTAAAA AGAATAGGAT 300

TTTGTATTAA TAAGAGTATC AAAAATTAGT TTACTAGAGG ATAAAAACAC TGTAAATGAA 360

ATACTTCATA CTAAGCTTAT ATTCTGTACA AGAATGTAAT ATTGTGTATC ATAAAAATAA 420

AAAATATAAT CCTAAAAACG AGCGATGGAA ATATTTATTT CTTTGTTTTA TGAGGTCAAA 480

AATATATAAA AAAGACTTTT TACTGAATTA GAGCGATTAA ATTTATTTGA TTTTTCAGCA 540

ATCCAAAAAA AATTGAAAAT TGGTTTAAGA TTTTAATAGA GATTTTAGTA ATTTAAAATA 600

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GCTCTTTGTT TCGCATATTA GTTTTGTCTA AATCAGTATA GAATCTATAT TTTATACCAT 960

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CAAGGTTGGC ATTAATCAGG ATAAAAAATT ATCATTTGGA TTTTTATTAA GTTCAATAAT 4380

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TTATCCCAAG TCTTACAGGT TTTGACAATA TTTCATTACC TCTAAGATTC TCAAGAGAAA 4800

GTGCTAAGCG ATTAAAGTCT AAAGCGGAAG AATTGATAGA ATTTTTCAAA CTTTCAAAGT 4860

TTGTGAATAA AAAACCTAGA TATATGTCTG GAGGACAAAG GCAAAGGATA GGAATAGCAA 4920

GAGCCTTTGT TTATGATCCC AAATTAATAA TTGGAGATGA AATAACTAGT CATTTGGACA 4980

AAGAAACAGC TATTTTTGTT TATACTTCAA TACAAAAGTA TCTTAAAGAG AAGAATGCAA 5040

TTGGGATTTT TGTTTCTCAT GATTATAATT TAAAAAATTT GGCTAATAAA CTTTATAGAA 5100

TAGAAGATGG AGTGCTGTCT TTAGTGGGGG GTGAGTGTGT TTAAATTGGC TTTTTACAAT 5160

ATCTTTAGAG ATTTAAGGCG TACAATCATA TTATCTTTAC TTCTAGCAAG TTCTGTGGTA 5220

TTTTTATTGG TTTTTGTTGG ATATATGAAC TTTAGTAGAG AGGGGATGGA AAAGAGCTTT 5280

GTTAGTTCAA GTGGCCATAT TCAAATTGCG AAAGAAAATT ATTTTAATCC TAAATTTAGC 5340

AACCTTAAGA ATGGGCTTTT ACTTGAAGAA AAGGATATTA ATTTGATACG GAATGAAATA 5400

GATAGTTATG ATGAATTACA ATCTACCAAT TTAATAGTTA ATTTTGATGG ACTTCTAGGC 5460

AATTCTTCGA CAAGTAACCC AGTTTTTGCA TTTGCCTATG AAGATCCAGA TATAATTACA 5520

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AGAGCAATTG GTTTAACCAA ATTGGAACTT TTTTACTCTC TATTTTTAGA AATTGTTATT 6120

ATTTCTGTTG TCAATATTGT TGTAGGAGTA ATATTGGCTT ATTTTGCTAA ACTTTTTATT 6180

CAGTTTCAAA AAATTAGCTT TACTCCTCCA GGCTATTCAG AAACATACTA CATCAACATA 6240

TTTTATTATG CTAGTGATAT AATATATGTT TCAATTTTCA TGTTAATTCT TGCTATTTTT 6300

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CAAGATTCCA ATTAAGGTAA GCTCATCCTA TAAAGTTAAG GGCGCATCTG AGCTTAAAGA 6720

TATTTTGGGT TTAAGTTTCA ATACAGATTT CGTTTTATTA AAATCCGAAG ACAATAGGGT 6780

TGAATTTCAA TCAAAAGAAA AATCAATATA TCCATTTGTA GATTTATTAA AAATTAATAA 6840

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TTTTAACGAT TCTAGTACCA AAATTTATGT GGAAAATATT ATTAGTACTA ATCTGAATAA 7020

CTCTATTTTT AGCTTAAAAG GATTTAATAG AATATTTGAT TTATATTCAA GGTATATAAA 7080

TTAAGAGCGT AGAGGGATAG TTAATATTGG TATTTTTTAA AAAATTTTGT TTAATTCTTT 7140

TTTTGGCTTT ACCTGGTTTT TTGTTTGCGG AATCCTCTTT GTTTTTAAAG GAGCATTTAG 7200

GATTCAAAAC AAAATTCTCT TTATTGTTGC CCGATGAAGA TAAAAAAGAT TTTTTTGGTA 7260

CAGGAGCTTT ACAATTTGAC AATGAAGCTT ATTTAGGATT TTTATTAAAC TATAAGATAT 7320

TAGAATTTGG CATTGCACCT TCTTTTATTG TGCAAAATAA TGATCAATAT TTTAGTTTTA 7380

ATAAATTATT CTTCAATTTA AGCTTTAATG ATTTTATTTT TAAATTAGGT AGGCAAAATT 7440

ATTATTTAGG AAATGGATTA ATTGAAAATA TTGTTTTAAA AAGGACCACA ATAGAACCAG 7500

AATGGTTTTT TGAGTTTTAT TATTTTATTT CTAATTATTC TGTCTCTTTG GGtTCTATGt 7560

TAGACAAArA AAgCTTArAT AAATTTTCAT CTCCCAAATA TTTATCCCCT TGGCTTTATT 7620

TTCA 7624

(2) INFORMATION FOR SEQ ID NO: 19:

(i) SEQUENCE CHARACTERISTICS:
(A) LENGTH: 7074 base pairs
(B) TYPE. nucleic acid
(C) STRANDEDNESS: double
(D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 19:

ATTGAGGCTT ACAATAAAAA TTCTGATAAC ATTAAAATAG ATAATAAAGC GCTAGAAACC 60

ACATGGAAAA AAAATACTCC CATCCTCTTT ATCTACTAGA TCAAGCTGAT TAATTTAAAC 120

AAAAATTAAT AAGTTACACT TAATATTTTT TAAAAGAGAA GTTAATTCTT CTCTTTTTTT 180

ATTyGTACAA TCTAAATCCA CGCTAAAAAC ATGCAATAGA AATTAAATTT TCAATATCAA 240

ATAAGTTTAA TATTACAATT TGATAwTAwT ATCAAATTGT AATATTATAA ATTTGAACTA 300

AAATTAATAA ATATTAATTC AAATTTATAA GGAGAATATT TTGAAAAACT TTAAATTAAA 360

TACTATTAAG CTTAACGTTA TTACAGCAAT ATTAACTTTA ATTTGCATAT CATGTGCACC 420

TTTTGGCAAT GTTAATCCAA ACAAGCTAAA AAATCCTATC ACCTCTAAAA ACCTAAAAAA 480

AmCAAAGCGA AGCAACCATT CTAGAAATCT AAAAAAAACA AACAGTCACA CCAATTCAGA 540

AAATTCAGCA GAAAATAACC AAAATCTTGA AAATGAATCT CAAAATTCAA AATCTTCAAA 600

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CACTTAATAT TTTTTAAAAG AGAAGTTAAT TCTTCTCTTT TTTTATTCGT ACAATCTAAA 1260

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TGCACAAGCT CTTATTAGAC TATCAAAATA ATACAAATCG TATACAAACA GATAAAAATG 2400

AACTTAAGTC TTATGCAGAC ACACTTTTCA ATCAAATGAC AAAAAAACCC GAAGAAGCAC 2460

TAAAGCTAAA AAATACCATA TGCTCAATAG AGGACCTTTA ATTTATATAT TGAAATTGTC 2520

ATAATATAAA GACCTATCTT TTCCTAAAAG ATAGGTCTAA AGTTGTTAAA TACTATTTCT 2580

AAAAAAAAGA TTATAAATAG CTTCTTTATT TATTCCGCGC TAGCTTTTCA GCTCTACAAG 2640

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CCATTCCTTT TGAATTTATT TTTAACTTTA GGAATAGGAT CCTTTGTCCA AGGAGATTAT 3360

ATTGGTGGTG GCGCAGTGCT TGGATCTCAG TTATTAGGAG GAATACTTTG CATAGCTGGA 3420

AATATTCTTG GCCATACAGA TGATGAAACA AGAGCAACAA CTGGGCATAT AATAACAACG 3480

ATAGGAGTAG GCACGATTAT AGCATCCCAC ATAGCCTCAC TTATTATTCC ATTTACATTT 3540

GCAAATAAAC ACAATGCAAA TCTTAAAAAA AGACTCGGCA TTGATATTGC GGGTTTTGAA 3600

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AGCTGTTCTC TATAAGGAAT TTTTTTTCAA CAATCATAAA AAGGGTCTTA TTAGATCATC 4620

AAAACAATGA AAATTCTATA AAACCAGATG ATTCTAAATC AGGAACCTAT TTCGATACGA 4680

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AATACTGATC AAAAAGGCAT TAAATACCCG CCTACCGAAA AATCAAAGCC CAAAACTGAA 5520

GACTCTAAGC AAAAAGAATT AAAGCCTAAA ACAGAAAAAG AACTAAAGAA AAAACAACAA 5580

CTAAAAAATA AACTACTTAA TGATTTAAAA AATTCAATAG AAACAGCTAA TAAGCATAAA 5640

GAAAAGTATA AAAAAAGAAT GAAAGAAGAA CCCGAAGATC AATACGGGGT ACAGGCTTTC 5700

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TAAATTATTA GAATAAATAG CAATTAAAAT TACTAAAAAT GTGATTACAG CAATAAATAT 7020

GTAAATATGT GAAACTATTT TTATTAGAAC TGACAATAAA GATATTATGG nTAA 7074

(2) INFORMATION FOR SEQ ID NO: 20:

(i) SEQUENCE CHARACTERISTICS:
(A) LENGTH: 6810 base pairs
(B) TYPE: nucleic acid
(C) STRANDEDNESS: double
(D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 20:

AATAAAAAAA CACATATTAT GCTCGCAATT TTCAAATAGT GTAATTAAGC TCTTGTATTT 60

AATAATAATC AAATACCTTT CTCTAAAGTA TAGTCAAATA ATAGCAACAC TAAAAAGTAG 120

TAATATTCAT TAATGTGAAT CTCTAAATCT CTGGCATACA TTTTAAATGT CATTCAACGC 180

CTTATCTTCA CAGGATTGCT ATTAATAGGA GAAAGTCTAA TGCACTCAAT TCTCCATTAT 240

TGCTATCCAC TTAAAGAGAT GGGGGATTTG AAAACATAAA CATAACTAAA CCATAATAAT 300

GATAAAGTTT TCAATCGAAT GGGTGAAAAA GTATAAGAAT ATTTAAGCTC TTCGAGATTT 360

TTAATTAAAT TATTATATTT GTCTTTTATA GGCCTTAAAT TACAAGATTT TACCTATGTA 420

ACATAGCCTC CTACAAACAA TTTCCAAACT TCTCCTTCTT TTATTTCGTT TAACCTAGAT 480

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ATAATTGACC CAATATTTAA AAAACCATTT ATTTCTGCGA TCTTACCTTC TTTGTCCCTA 600

TACTTATCTT TACACTCTCT TAACTTTTTC GATGAAATCA AAACCCCTCC TTATATTTTT 660

TTATATATAA AAGATAAAAC AAGAAAATAA ATAAACCAAA TATAGTCTTT TTGGCTTGTA 720

AAATTTTATT GGAATTTCAC TTAATATTTC TTTTATAAAT TAAATAAAAA CATTGATTTT 780

TGCACAAGCA AGGGTAAAAT AAATTCAAGT CTTAATTAAA GATTCTTTTT TTAATAAAAT 840

CTTTAATCAA GACTTGAAGT CTTTACAATT CACCACAATA ATAACTGGAA GATCTAAACA 900

AATAGAACCA CCATTTATAA GTAATATACA TCAAATTTCA AATAAGATCA ACCCTTATTC 960

ACAAATTTTT TATTTCTATT ACTAACTACT ATTTTTTAAA TCAAAGATTA TAAAAATAGT 1020

AGTTTATATC TTATGTTTTA TATACCAAAT TATTCATAAG TGCTCAACTT TTAAATTCTT 1080

CCATAAACTT TTCTAGCAAA TCTTTTTTCT CTATAAAAAC TCTATCTAAA AAGTAACCAG 1140

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TAAATCACTC ATTATCCTTG TATGATATAT TTTTCTGTTA TTTTTATCTT CCACTTTGGC 2880

AAAGATAAGT TTTTTTCGTG TTTTTTTTAT TTCTGATTTT TTGATTTTGA GACGCTTTAA 2940

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TAACTTTTCT ATGGATTTAC GGCTGTTTCT TTTTTCTTCT TTATTATTAG ATATATTATA 3660

AAAACACTCC CTATTTTTTA CACTACTATT TTCAATGCAA TTCTTATTAA AATATGAATC 3720

AACACGGCTT TTATATTTTT TCTTTTTTAT CTCTTTAAAA TGTTTATTAA TTAAATGGTA 3780

ACATACTTTT TTGGGATATT TTAGTTTGTA ATGAACTTCG GTCCCCATGT TAATACCTAA 3840

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GCATAGGGGT TATTCTTACT AACTCAGACT GCACAACTCA CTAGAGGACT CAAAATTCTA 4380

TCCAAATATA ATAATCTTTA GAATTTTTAA GTCTCGGTGA AAAATTGGGT TTTTTGAAAC 4440

AAATCTTGCA TCACTAGAAG AAAGTTTAAT CTCTTCAGTA GATCTGGCTG ATTTTACTAA 4500

TACAATTCTT TTAAGTATTT TATCTAAAAT TTTCTGTGTC TTTTTCAAAT CACTTGCTAT 4560

TCATCATCTA ATTCACTTAA TTTTTCTTTT CTCATTTTGA TTTTGGCGTC AAAACTCTCT 4620

TCAAGTCTGG GAGTTGCATG GTCCTTCAAA ATAATCAACG CCCACCCCTC CATGACTAAA 4680

ATCTCTACAC ATATCGGTTA TTTTATCTTG CATAGACTTG TGGACTTCAA CAAAAGATTG 4740

ATCTAGAGCC ATTTATCAAA TTCATTAACT TAAATCAATA AATTTAAAGT TTATGTAAAA 4800

CATGCTAATT ACTATTTCTA AATCAAAAGT CATAATTTTT TATGACAGCT TTTTAAGAGT 4860

AGTCATCATT AAACAACTTT ATTATTAGAG CTTAATTAAG CATTTACACG AGGCAAAGAC 4920

TTAATTAAGC TCTAATGGCG GTCACAAATT AAGGTAAATT TTAAATGAAA AACTTATCCT 4980

TTACTTAGAA GGATACACCA ATTTAAAAAT AAAAGCAATA AATCAACAAT TCAACAATCA 5040

TATTAATAAA CTATTTAAAT TTTTTCTTTA AATTCTCTTA TAATTTGTTC AACCCAATCT 5100

TTTTTATCGA AAAAACTTTG TCTAAAAATA ACTCTGTAAA TTTACAGTTT TTTATAAAAT 5160

AAATAACTTT CTTGATTTTT AAGCTTAAGT CTTTTATATC CATCGAGTTT ATAATAAACT 5220

TTATTTTTGG TTTCCATTAA ATATATATGG CCTCTATCTT ATTATAGATG TTCCCTTTTA 5280

CATTATCTTT TAATTTATTT TTTAATAATT CAAATCTTTC TTTATCATTA AATGAATTTA 5340

TATCATTATT GACATCCAAA ATATAATTTT TTATATCTTT TAGTTTATTT TTTAATAATT 5400

CAAATCTTTC TTTATCATTA AATGAATTTA TATCATTATT GACATCCAAA ATATAATTTT 5460

TTATGATTTA AATTTCTTAG ATCAATAGTT ATTTCATTAA TATTTTTTAT TCATAAAATC 5520

TCCTCGTTTA GTTTAATTTT ATTACTTTAT AAATTGTCTG GCGCTGGACA GTTTATATTG 5580

GGGGTTTCCC ATTTTTTTGA ATAAGTTCTT ATATAAAATA TCAAATTATT TAATATATAA 5640

TATTATTCCA ATATGGAATT TGATTTAAAA GCAAATACCA TATGTTTGCA ATTTTTTTTA 5700

TTTGGTTCCA TTTTTCTGAT ACAATGCCTA AAAAGCATTC TTTCTTTGCA TTATTTTCAA 5760

TAATTTTTTG TGTGTATCGG GCTTTTAAGC CTAATGTGTG GCATTTAAAG AATATCTCAA 5820

TACTATATAG TCACAATTGA CTTTTAATTT TGTTATTTCT ATTTTCAATT CTGAATCAAT 5880

AGCTTTGTGC TTATTTAGGT AATAGAAATA TCTATTTAAG CTGTATAATG TCAAATGGCT 5940

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GCTGGTAATT GTTATTATTT TTGTTCTTTT TCTATCCATT TTGTTATAAG CCCTCCATTT 6060

GATAATTTTT TGTCGTAAAA TTCATATATT TCTTTTTCCA AAGTTAAAAT AATTTTGATA 6120

AGAGATTTAT AATATTTTTT TTAATTTTAT CTTTTCTTAA TAAATAAGAA ATTCCTACAA 6180

TATAACAAAA TATGCTTCCT TTCTTAAATT TAAATTGTAT GTAGTGTATG CTTCTGTTTT 6240

TTCCCTATCA AACAATTTTC TAAATTGGAT AAAAAATCTG TGTTTTTGAT TCTTTTTTGT 6300

TTCAAATGCA TGAAAGTCTA AAAGTATTTT CGTATGATAT ATCTTTTTTT CGATTTTAAC 6360

AAAAATTTTG TTTTTGTCTG TTTCGATTTT TTTGTATTTT TCTTTAAGTT TTTCTAAAAG 6420

TTTTTTCATA GTCTACCTTC TATCTAATAA AGTCTACCTT CTATCTAATA ACTTTTCGGT 6480

TGTCAAAAAA TTTATTTGTG TTTTTTATTA AATTTAGTAG ATCTAAATAG TATTTGTTGT 6540

TAAGTAGATA GGAATATTTC GGTTTTTCTA TTTGAATTAA ATACTAACAA CACAAAAAAA 6600

TATAAAAACC AATATACTTA ACATATTATT TGAACAATTA AAATCAAAAT TTGAAATTGA 6660

CAATGAAATT TTACTAAGAA TGGTAAAAAA GTATATTAGG TTTATCTATG TTTTTTTATT 6720

ATTTGGTTTA AATCATCATA TTTGTTGTGA TCGATTATAA AATGAGGGCC TATGTTTGTA 6780

TTGGTAAATA TACCATTTTA AAAGnCCTTT 6810

(2) INFORMATION FOR SEQ ID NO: 21:

(i) SEQUENCE CHARACTERISTICS:
(A) LENGTH: 5805 base pairs
(B) TYPE: nucleic acid
(C) STRANDEDNESS: double
(D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 21:

CTTGCATGCC TGCAGGTCGA CTCTAGAGGA TCCCCGTGTA AAAATTGAAT CAGTGCTTGC 60

TATCCACTCA CCTAGCAAAA CTCTTGCTTT ATATGATGGT ATATCTTTAT ATAGCTTTTC 120

TTGTGTTTCG ACAAATCCTT TACTAAGTAG CACATTATCA TAAGTTGTAA AATTATATGT 180

CTTAAAGGTC GCTATATTAT CAATATAATC GGTTTTAAAA TAGTGTTCTG GATGATCrGG 240

ATTAGTATCA AAAATAATAG TTTCTTGCCC GCATCTTAGT CTTTTTAAGA CyTCCTCTAA 300

AGTTTGCTTG TGTAAAGTTG TAGCCTCATT AACAAAAATA AGTGCCGAAT TACTTCCCCT 360

AAATCTTTCA AAATCACTTG CCTTATCTCC wCCATATAGA TTAATACGTA GTGAATCAAT 420

CAGAATATAT GAATTATTTG TATGTCTTGG AATATAAGGA ATTTTAAGAA GTTTACATAG 480

CTTTTCAAAT TGyCCCAAAA CATTAACTTC AACTGArCGT TGTGAATTCC CAATAATAAA 540

ATTATTArTA TCACyAGAGT ATAACTTTTT ATTTTCAATT AAACTTTTGA GAAAAAGATA 600

ACATGCAAGA TACGTTTTGC CGCTAGCTAT GCCGCCGCTG AGTATAATCT TCTTTTCATT 660

ATTTTTTTGA ATACTTTTTA TAACATTTTT TTGTTTTATG GTTAACTGTT CTTTTTCAAA 720

CTTATCAAAA TTAATTGAAG AATTTGTTAG CTTTACAAAT TGTGATATGT CAACTCCATA 780

TTTATTTTTG TATTCCTTTT GTAGTGTTGT AAAAAGTTTT GTTTGATATA AGTTCACTTG 840

TACCCTTTAT AACTTTTGTA ATTGCCTTTA TTCTCAACTT CATGAGCAGA AGAAAGTTTT 900

TTCATACATT CATAGTAAAG CTCCATTTCT CTTATTGAGC ACTCCTTTAC ATATTCATCA 960

AGCTCACTTT TTAAAGAATT AATTTCTCCA TTAACAACTT GCTTGTTTTT TTTACTACTT 1020

GCTTTATTTA AAGCATCAAT TTCGGCTCTT AAATTTTCTA TTTTAGTATG CATACTAAAA 1080

AGCTCAACAC TAGAATATCG CTTAAATGCA TGTATAAACC CTAATTCTAA ATTGGCTCGC 1140

TCTAAATCCA ATTCGCTTCT AACTTTCCTA GCGTTAACTT CTGATCTAAA GGTTTGCGAC 1200

AAAAGGTGTT CAAAAGTATC TTCACTAATT GTTAGTCTAG AGTCCTCGCT AACAGAAGTT 1260

TCTCCACTTT CCCACTTTTG TCTCATTCTC CACACATTTA CTTTAGAAAC CCTTAGTTTA 1320

GCTGCTATTT CTTTATCACT TAACGATCCT TCTCTAAAGT ATGCAACATA ATCATTAAAA 1380

GACCTTTTGG CTCTTTTCAA GAAAATTCTC CTAAAATAAC AAAATTAACA ACTTATTACT 1440

CTAAATAGTA ATTCAATTTA TTAATTATTA ACATTAACTA TTATCTTATT GATATCTATT 1500

GACAGATGTT TGATATTTAT TGACTTTTAT TGATTTAGAA ATAGCGATTA ACTAATTTAT 1560

TAAATTTTGC TACAACTTGA CTATACCAAT TTGGAAAAAT CTTTTATTGT TTTTATAAGA 1620

TACTTCCTTG TGCAAATTCC CTTATCATAG TAGTGCATGA TTATAAATAA TATATCTACA 1680

AAGTTAAGAA CCCCGTCTAT TGATTCTCGT TTTGGATCTT TTCTACCAAT CTCATCCATA 1740

ATATCAGAAA CAATACTCAA AAACCGGTCG GAATATTTCT CTCCCATTTT TTTCAAACAT 1800

TTTTCTAGCA TTCCTTTCTT ATAGCTACAA TCAAACTTCA TATTTTGGAA ATTTTTATAC 1860

TTTTCTATTA TATGCTTATC TTTCAGAACA TCTTTAATAA AATTTTTTGC AGGGTTTATA 1920

ATACCATTTA TAATTTCATC AATACTTTTT CCGGTTTTTC TTTTGTATTC TTGTATTGAT 1980

ACAAAACCTC TATAAAAATC CTTTTTTTGT TCCAGAGTTA ATTGCTTAAC TCCATAAAGT 2040

TTGCTTATTT CCTTTTTTGT GCCTGGGCTC AAACAATCAT AAAAATCTGA AATTTCATTT 2100

ATATACATCA AAACTCTAGA GAGTATATAA ATAAACAATA TTTTTAACAA TAGTTTTTGG 2160

CATTTTCACT TCATAAGTGA TATAGCTTAA ATAGAAAAAC AAGACTATCA ATTGGCCTTG 2220

TTTTTCTAAA TTCGATTAAC AATAAAAACT TTTTCTTTAT TCTTTTTTAC CTTCTACAGT 2280

TTCTCCAATA TTTTTAAGTT CTTCTTCAAT CTTTTTAAGC GAATTAGTTA TAACTTCATT 2340

GGTCATATCG CTAGTATTAT TACCACTAGT CATATTTTTA AACCCCAAAC TATTAGCACA 2400

CTGCCAAGCT TGTAATCCAA CACCACCTTG TCTTTGGACT TGAGAACCAT GCGTTACTCC 2460

AGTTGCAGAT TCAGCTTGGT TTTTATACTC TTCAAATTTC TTTTTAGCCT CTTGTAAAGC 2520

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AAGTCCTTTT CCGTTTAATT TTTCTTTTAT TTTGTTTATT TTTTCTTCAT ACCCAGAATA 2640

TGTTTCAATA GAAGTTTTTT TAGAGTCTAA TTTTTCTATC TTATTCTTTA ATTCTTGAAT 2700

TTGTTTTTCA ATTTCTTTTG TATCTAATGT GTTTAAATCT TTTGTCTCTA AAATTTCTAA 2760

AAATCCTTCA ACTTGTTTTT TTATCTCTTG TTCTGTTTTT TTTACATTTT GTTCTGAACT 2820

TTCTAAATCT TTCACCGCCC CTTCTAAATC TTTACTAGTT ACATCATTCT TACAAGAACT 2880

TATCAGCATA AAAATAGCAC AAATAATAAA CATTTTTATT TTTTTATTCA TAAAGATTGT 2940

CTCCTATGTT GTAAGCGTTT ATTTCAAATT CTTTTTAAAT TCATCTTAAA TTCCTCTAAT 3000

AAAATTGCAT ATGTATTATC TCTTCCTAAA AATGTAATAA ATTGAGAGTC ATTAATAGTT 3060

TCTATTTTTG TTTCTATGTT TTTGATTTTA TAATCTTCTC CTTCTTGATA ACCATAATAA 3120

TCCGCATTAA ATGCTCCTCC TCTAATCATT GAATTTACAA AGTTGTTAAT TTCTGACTCT 3180

TTTACTGAAA ATAATCTAGC CGTGTGCCCC CCAACATTTA AATCTGTTAC AATACTATAT 3240

TCTTCTTTTA TTAGATCTAA AGTTCCTAAA TCTATCCAAT TGCCGATATT TGTTTTTTTA 3300

TATTTAATAT ATACATTAAA ATCAAAAAAT TCGTCAGTTT TTATGCGAAG ATATCCAACT 3360

AAAGCGCACC TTCTGAAATA AGTAATAGCT GCTCCAACCA TTTGGGGTAC GGTCTTTCCA 3420

TTTTTAGAGC TGTTTTCATT GGTCCATTGT AATTTTTCTG TAAAAGTTGG CGTATCAAAT 3480

GAGTATTTGT ACACAGTGCT TGTACTGTAG AATGTCGTCC TAATATAATC AACTATGCCA 3540

TATTGCCCCT CTACAGAAAT TGGATCTTGC GTAAAAAAAA GATCCAAATT ATGCTTATCA 3600

ATAACGTTTT TAATTTCTCT AACTATGTCA TCTAAACTTC GATAATTATA CTCGTACCCC 3660

TTAGAATTCT TATTAATCCC TGATAAATTC ATTCTTAGGG TTTTCATATC TTTTCTGAAG 3720

CTTATTTTTG CTTGAATATT ATTTTGCATT TCTTGATTTT TTTCTGAAAC ATTGTTCATG 3780

GGGTTCCCCT TCTAATTATT TTTCAAAATT TTACCTAGCC AATACTTTGC TCATCATATA 3840

AAGATAAAGT ATTATTTTTA CAAGAAATTA TCAAAACAAA AACTGCACAA ATAATTAAAT 3900

TTTTCATTTT CTTATTCATA AGTTACTCCA TAAGTCCCAA TATTACCACA ACAGCTAATA 3960

ATTGCAATAT TTCAAAGATT TAAATATATA ATTTTGTTAC ATTTCGAATT ACATTGTTAC 4020

AAAACTAAAA TGTAATTTTA ATCAAACTCA TAAAATCTCT CCATTGCAAA TGCACTACTC 4080

ATTATAAGAG ACTACAAAAC ACATACAAAT TAAATTTCAA AGTCTTTGCT ATATATTACA 4140

CAAAGTATTG TATCTTTCTT GTGTACCACC CTTAAAAATT GCCTCTTCTG TTTATCCCAC 4200

CTGCTCTACA GCCCAGATTT TGCATGCAAT GAGAACTCAC AAAATTTGAC TAAAATTTTT 4260

AGTTTTTGGT ACAATATAAA TTACATTTTT CATCTATTTT TATTACTTTT ACTTAATTTA 4320

AAAGTAACAC TTATAAGGAG CTTGTCTTAT GGATACTAAT AATTCTTTTA ATTTAAATAA 4380

TTTCAATATG GATTTTACGC TCAAACTATT TCAAGAATAC CAAAATGCAT TAAATAAAAA 4440

CAAAATTCTA GAAAATGAAA ATAAAATTCT TAAATCTCTA GAAAATTCAC CTAAACGTAA 4500

GAAAAAAAAT TCAAAACCAA CTCCTAAGTT TTATTTGACC CCTAAAAGTA TTAAATTAAT 4560

TCTAAAATGT GCCAAAACCC TAAAACAAAT TGACCCAATT TCTGGTTGGT TTGTGCATCT 4620

ACTCTTAATA AGTGGATGTA GAGGCACTGA AATGCAAAAA GTAAAAATGC AAGATATTTC 4680

AACTTTTTTA AGCAAAACCG GAAAAACTTT ATATACTATT AAAGTAAATG TGGCAAAAAA 4740

AAGAAATACC TCTTGTATTA GAGAAATTGT CATCAACTCA GAAGAGTTCG AGGCTATCCA 4800

AACAGCACAT AAAAATCATT TCCAAGAAAA AACTCTTGAC TCGCGACGTA CTTATCTTTT 4860

TCAAAAGAAC AAACATAAGT TTAAAGATAA CCAAATTGAT ATTGTCCATA TTTCTAAAAA 4920

ATTCAAAAAT CTTCTTAAAA AATCGGGATT TCGTGTAAAT AAATCTCTCC ATCTATGTCG 4980

AAATTTATTT ATTTCAAATT TGAAATCTAA CGGCTACAAT TCTTTTCAAA TTAAAGAACT 5040

TATGAAATAT TCTTCAACCA ATGAAATTGA TAATATCTAC GGmCTmTCTT CTGCTAATAA 5100

AATTCAAGCT TATGAATGTG CTAAAAAGTG CCTTAAACTT TAGTAAAACT ATTTCAGTTT 5160

AAATATACGC TTTGAAGTTA CTTTAAATAT TTTTCCACGw GGctTTcAAG TCAAGTGAAT 5220

CATATAATAT TTCTTTATTT TTTGTTGCTA TAAAGTGATA TCCATTAACC TTATCGATTT 5280

TAACTTCrCT TATTTCAAAT TCATTGGCTG CACCTAAATA ATTTAAAGAy TCATATCTCA 5340

CrCTACTTCT AATTCCGTAr TAATTAAGTA TCATACATrG ATTTATAATA AAACAATTGC 5400

TCTTAATATA TCCAAGTCCT ATAAATCTAT AATACGCTGC ATTTATCTCA TAmrCAyTAA 5460

ATTCACGTTT CTTAAATAGA CTTGTATAAT AATGTAGACA CAAAAAGTAA CAACCCCATT 5520

TTTGTATCTC CAGTCTTAAA GTTCTATTGT CTtGTTTTAT TTTATTAATA AGCATTAATT 5580

AATCCTCCTA TATTAAATTA TTTTTTTAAT TTTTTAGTGC ACCCATCACA TGGGGACACT 5640

AACACTATTT GCAAATGCTA TATTACTAGG GAAAAATACT GTACCACTTG TTGGTAGTAG 5700

TCCCTTTAAT CTATCTGATA TGCCCnTTTC TTAGGGnCAA GAGTAAACAC AAACTCTCCA 5760

AATTTATAAT TAACnGCTCA ATAGTGGGTA nGTATCGGAA AATGn 5805

(2) INFORMATION FOR SEQ ID NO: 22:

(i) SEQUENCE CHARACTERISTICS:
(A) LENGTH: 5377 base pairs
(B) TYPE: nucleic acid
(C) STRANDEDNESS: double
(D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 22:

CTCAAGATTA AGTAAAGCAT TGCGAACTGC ACTGTAAGTT GTGCTTTTTG AAGAAATATG 60

TTCTTCAATA GCAGCCCAAT AAGTACCACC AGGTTTCATT TTAGAAAAAA AGAGATTGAG 120

CTCATTAATT ATTTCTTCTT CAATTAATGC CAAAGAAGAC GCAATAATGT TATAAATTGA 180

GCTTTGATTG TCGTCAATAT TAaTmCCaTA ATTTACaCGC AAaTATTCTC TTTTmGmTCT 240

TACAaTATCC yTAaTTGTAC GTTTTAAAAT GCCAAAATCA GAATCAAAAA CTATrCTCAT 300

AAATTAAACT CCATATTyAA AACaTCGCCg aGAAAAAAAA GGATATATGT GCTTTGTTGT 360

CTTGTATAGT AGTTGAAATA TTTATTAAAT CTAAGTTAAG CTCTTTAGAT ATTTCATGAA 420

AATAATTTTT TACAGCGTGA AGATTGTTAA TTTTTAACAG CTTTAAAAGC AAGAAATAGT 480

CCAATCCCCA ATGAGGAGCA TAGCTTAAAC TACCCCTTAA GGTTTTTAAA AATATCAAAA 540

ATCTTTGTTT TTGTTCATCA ATTCCATCAA CAAGTGATAA ATCGTTATTA AATACCAATT 600

CAAAATTATT GCCTAATCTT AAATCCATAC TAGTAATTAT ACCATATTTA GCAAAAATTA 660

CTTAAACAAA CTATTAATAT CAGAATTAAT TTGAGCGGTT GCTATTTTTA AGCTTGATTC 720

GTCAATTACT GAGTCCCCTA TAATTTTTAT ACCATTGATA GCACTAACAA TATTATCTAG 780

AATTTTTTTT AAGCTAGTTG TTTGGTTTGC TATTTCAATT TTATTATTCG CTCTAATTTT 840

AACAGTATCA GAGATTAGAT TTAAAGTCTT TGGGCTAATT GCACTAAGTA TATAAAAATG 900

ATGTTTGTCA AAGTGAATAT CGTTATTTTT ATCAAAAATA TTAATGCTTG ATTGAAGTAG 960

TAAAACGywA TCACCTTTTG ATArTTCTAA ACTGATATTA GAGATATTTT TTGTGTGAAT 1020

TTCTAAATCT TCAAATTCGG GTATTGTAAC AATAGCyTCT TGAGTTTGwT GTTTAAACTC 1080

CTTTACAGTG CCAATTTTAA TTATAAAAAT GTTTGAATAA ATCCAATTTT TAACGTCCTC 1140

TTGTGCTAAA GCATGACCAT AAAGACGCTG GTTCATTCTG TAAATTTTAT AGTCTTTAGT 1200

CATTTTAATC CCCTTTACTT TATGCATTTA ATCAGAATCA TCAGAATCAT CCTTTAATTT 1260

AAGTATTAAT GAACACTCTC CTGCGTTGCT AAGTCTTGCA CTTGTTTCTT TTATGGTGGT 1320

TTTTATCATT TTTCCTAATC CATCAATAAA TGAAACTTTA TCGCCAATTT TTAATTTATG 1380

TGTGTAAAAT GTTTGTGCGT TCCAATATAC ACGTTTTATC TTGTATTCAC CTTCAATAGC 1440

AATTTCTTGT TGTGGAATGA ATTCAAGTCC ATAATCTTCT AAAGCTTTGT AATTTTCATC 1500

TGCTTCAGTT GTTATATCAT TAGTAAATAT AAAATTACAT TCAACATCAA AGCTGTCACC 1560

ACCAACATCA GCTATTACGT CATGAATATA TATTCCTTTT ATTTTGTCAA CAAATTCTTT 1620

TGGTGTTGTG GCATAAATAT CTTTTTCAAT GATTTTAAGA CGATCTTCTT TGTCCATATT 1680

AATAATATTG CGATTAGGAA ACACAGATTC TATTGCCTCT TGCACCGTTT TCCCCTTGAA 1740

GTTTTTGTTC TCTAATTTTC GATCAAAGAA GTTGCTACTA ATTGCTAACC GAACGTCAAG 1800

CTCAACACTA AAATCACCAC CAGGATAATC AGTGCTCATA GGAGTTCCTA AATACCCCGA 1860

CATTATGAAA TCAAAATTTT TTTCATGAGC AAATTTTTTA TAGTGTATTT TTACTATATC 1920

TCCTACTTCA ATGTCGTTGG TGAAGTCTAA GGGCAAATTC CAAAGTACAA TTTTTGTTCG 1980

TTTAGATTTT ACAAAATTAT AATTTGAAAA CTCATTAGAT ATGGTAATAT CAATATGAAT 2040

TCCATGTTGT GTATTTATTA TAATTTTGGG AGTTTCTTCA ATTAAAGAAT CTCCAGAGCT 2100

TGTATCCTTT TCTAAGGATT GTTTTGCTTT GTAAAATTCA ATTTTAAAAT CATATTGTAG 2160

TAGCAACATT AATTATCCTT TATATTTTTC CAATTCAAAT GTTTTTACAA TTTCTATTGA 2220

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TCTATTTGCT AGTTGTTCTT TTGCAAGATG CACGTCAAAA CGTAGCATAG TATTGCCAAA 2400

AACAGTCATT TTGAGCAAAG AAAGCATTTC TTTATATAGT GAGGTTAATA CCCCACCATT 2460

TAATGAGATA TTTTCACCAA TCATTACTGG ATTGTAGCTT ACATATTCCG CTTTTCTATC 2520

GTAATAGTTG ATAACTGGTC TTTTGGAACA ACTGGTATTG TAAGTGCGTG CTATGAGTTC 2580

GGTTTTTGGT TTTATAAAAA ACAATTGAGG AACATATCCA AAACCTTTTA AATCCATTCT 2640

TGGAAATAAC ACTAAAAAAT TATCTGCTCC GAAAAGAGCA AATATTTGGG TTATTACATC 2700

TCTTATTATT CGAGTAATCT CCCAGAATTC TTTCTTTTCA ACATCTTTGA TTTTTTCCTC 2760

GAATTTTTTC TTTTCAAGAT TATTATTGAT AATATTTTTA TTGTTAATAT CCATAGAATT 2820

TCCTTTTTAT TTGAAATTTC CTAAATCATC ACCCCCGGTG TTATTGTTGT TCTTTCCAAA 2880

AATCGTATTA TTAATCCAAC CAGTCGCGAA GTTATAAGTA TTACCAACAA CATTACTAGC 2940

GGTATTATAT AATGAGCTGC CAATGTTTTT TATTCCGTCA AAAGCACCCT TTATACCGCC 3000

TGTTATGGCA TCAACGATTT GTTTTAAGAT TTGTTCAAAA TTAAGGTTCA TAATCTTTTC 3060

AAGAATACCA GAAATTTTGC CAATTACTGG CATCAAAACA TTATTAGTTA CGTCTTCAAG 3120

ATTTTGCTCT ATTTTAGCTA AATTACTTTT AGCTAAAGCA ACTTCATCAG AATGACCCAT 3180

TATATTCAAA GATGATATAT CTTTTATTAA GTCTTTAATT ATCTCTGGCC TTAGCTTTTT 3240

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TCCAAATTTG TCTATAGGTT TCAATATATT GAAAGCTTCT GTAAGTTCCC CTTTAAGTAT 3360

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AGCTGCTAGT TTTGTTGCAT TTAATACATT CTCTCCCTCT TCATCATTAA TTTTTAATTC 3480

CCTGATGTGG CCTTTAAGCA CACTCGATGT TCGCAGAAAG TCTTGTTGTT CTAATTTCCG 3540

CTCAAATCCT TTCATTCCCT TAATAGCACC CAAAATATTT TCACGTTCTT TTTCACTGTA 3600

AAACACTTTA TTAAGTTGTT TTAGTTGTTC TTCTTTAGAT TTATTTTCAA CTGCTTTTTt 3660

AGCAAATCCA TACATAAATC CAATAAGCCC GCCACCAACT TTACTTACAG CGTTGCCAAT 3720

GACATTTCCT AAAGCACTAC CTATTGCAAT TTTGGCAACA AGTCCTTTTC CTTGAGAAGC 3780

TGCGAGCATT TTACTTTTTG CTTTTGATTC TTTTGCAAGC TCTTTATACT CAAGGCGTCT 3840

TTTGTCTCTA TCAGACATCA AAGATCTTTT GAAAGCTTCT TTTCTTGCTT TCTCAAACCC 3900

CATGCCCTGT TTCATAAGCT TTTTAGTCTG TGTAAGTCTA TATTTCTCAA CACGCTCTCT 3960

TAAGCTTTCA AATTTAGATT GTTTGCTAAG TTCTTTTTTC TTGTCCGACA AATTATTTTT 4020

TACAATATCT TTAGTGCTAC CCAAACTAGA TTTTTTGGGT TTAAGATATT TTTCCATTTT 4080

GGAAATATCT TGTTCAATAG CCTTTTTTGT TGCAGCATGA TCAAGAATCC CTTTAAATTT 4140

AATGGTGAAT TTGTCGCTCA TTAAGCCCTC ACTTGCTTAA AATTAATTCA TACAATTCTT 4200

TTTCTAATTT AATATCAGCA AGTCTATTTA CCTCTAAAAG CTCGTCATAA GGCAATTTCT 4260

TTACCGAGTC GTATGAGCAA ATATTCATAA TTACTGGAAA ATAGTATTTG TCGTTCTTAA 4320

TTTCGTCAAG CAAGTTAAAA TACTTTTTTC TAGTCTCATT AAGACTTGCA ATGGCTTTAT 4380

CAATATCTCT ATTTCTTTTG ATCATTTAGC AACCAGCTCA TTGGAATTTG ATGTACTTGA 4440

TGAAAGCGAA GTGGCTACTT TTTCATAATC AAAGTTTTCA TTGATATAGT CAAAAGTAAC 4500

AAAGTCCCCA ACATTATTTT CATACTCACT CAAATATACC AAAGCGGGTT TTTTTAGATC 4560

ATTGTCTAAA TGAAAAGTAT TAAATTGTGC GGTGTAAATT ATTGCAACAA GATAGTCTTT 4620

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AAGACTCTCC TTTTAAGAGT TGTTTGGTTT AGATTTTTGG CAATTAATAG CCCTAATTTC 4920

AAAAGATACT TTTTCGGCCT CAGCAGAATA ACTTCTTGAA GGCTCTTCAG TAAAAATTGC 4980

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CTCTTTTCTC ACATCCATGT TGTAAAACTG TTCATCAGAA AGTTCAGTTA ACAAAATGTA 5100

GTCATGACTA CCCAAAGTAA CTTCAATGTT GAAAACATAA GTTATCGTTT TGGGATCTCT 5160

TAAGCTTATT ACAGGCATAC CTTTATCTTC ACTACTAATC ACTGCTCTTG TTGTGGGTTC 5220

GCTTGTAAGT TCTAGTTTGC CACTATGTAG CTGTGTACCA CCAATTGAAA AATAAACTTC 5280

TCTTAAATCA TAAAATTGCA TTTTTAGCCC TCCTTTTAAG CACTTAAGCT GTTTTGATAA 5340

TCAACTATAT CTTGAGTAGa ATTACTAAAG AAACAGC 5377

(2) INFORMATION FOR SEQ ID NO: 23:

(i) SEQUENCE CHARACTERISTICS:
(A) LENGTH: 5121 base pairs
(B) TYPE: nucleic acid
(C) STRANDEDNESS: double
(D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 23:

CnTTTGCTGT TATAGTATCT AGATTATCtT CTGCCGAATT TATTACAGCT GTTAATGAAT 60

TTAAGGATGC TACTGAAAAG TATGCTAATG GCAATAAAGG AGACCATGCT GTTGATGTTG 120

TTGTGGGTGC TATTGCAGGC ATAGCTTTTG ATAATGAGAA TAGATTTGAA AGAGCCAAAA 180

TGTTTGCTAA TAAGGAAAAA GGTGCGGAGG TAGACAAAAT GATTGCTGCC ATTGAGAAGC 240

TAAGAGCTAC TTATACTGCA GTTAAGCCTA AAAATAAAGA TAAATAAATA ATAAATGTAA 300

TAACTGGGAA AATACAGAGA ATATTCTTAA GCATTTTCTA TTGAGCTTTT TGTATACATA 360

ATTTATTTTG GTTATGTTTT ATTATGTTTT ATTTTTTATA GGATATAAAT AGACATATAA 420

GTCTATAAAA ATATTTGTCA CACAAGACAT TCATGTAATC ATTTATTTCT ATTATTATTT 480

GATAAATTTG CTCAATATGG CTTTTTTAAA AAAATAGGGC ATATCTTTAA AACCCTTATC 540

AATAAAGCAG CATTAATGAT AATTTTGGAA ACATTTGCTA CACGTACAAG ATATTAAACC 600

ATTTTTTTTC TTAGCATTAA TATCATTATT ATATATTTAA AAATAAGCCA CGATTAAGGA 660

TCCAAATCCT CCTGGGACAA CATATGGGGT TGATCGTATA TATTTTAGCC CCTATTCCAT 720

TAAAAATAAC ATTTGATAAT TCATTTAATT TAAAAATGAA TTATCAATTA CAACAAACTC 780

CATGTGATAA GATGGATAAT AACTATCGGA TTTTTCTTAA TCCCATTATC TAGCATCCGG 840

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TGTTAAAACT AATGAAAAAT CCATTAATTT TAAGGAAGGT TTACAAAACA TAAAAGATTT 1080

CTTCCTGCAA AGTTATTTAC CTAAATATGC TTAATAGTAA TAGCTTAAAA GCGGATATAA 1140

ATATTATTGT AAGAAGAGAG TCGACAATTT TGATACTACT TAAAGGTTTG GATTATTCTA 1200

TTTCTGCTTT GTTTGTTTTA AAGCGATTTA TGGTTTGAGG CTTAATGTTG ATTTTTTTAC 1260

CTTTTTTTAT TTAAATTTAT TTACAATAAA GCTTTTACTA TAAATGAATT ATGAATATTT 1320

TAATAAGAAC AAGTCAkATC TCATAATTAG TGCAATCAAG AAAAAAAaTT AGCACATATC 1380

CATTTATGAT TTAATTTCTA CTATTTAGAT CAGTGGTGAT GCATTATTTT AGCTTAAAGG 1440

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ATCAAAATTA TCATTATCGC TATCATTTAT TTTTGGCATC ACAGCTTCTA ATATCTTrTC 1920

TAACAAGCAT ATTCTACCTA TCCCGATCTT CATATTGTCT GAAAGAGGAT ATGAGGCATC 1980

ATATCCTGAA GCTATCAATA GCTTTAAAGA TTCATCAATC TTTTTGGCTG ATTCTCTAAT 2040

ATCTCTTACA AGTGAGGGAA CCTCTCCCAA TGCTATACTA GCATTCCCCT TTTTACTACT 2100

TAAAACCTTA TCTTTAGATT CCTTGGCAGA AATACTAGCC TTTTCAACCA TATCTTTAGG 2160

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AAATTATTTG TTATATTTAT ACTCTTTGTT AGCTTTATGT TTTTAAGTTT TTAAACAATA 2520

TCTATCTTTT TGCCyTATAA TTTTATCTGC GCCCAGCTTC ATAATACACT CATCTTGAAT 2580

ATAACATGAA AAAATATAAT ATrCCAACTA GAAATCTTAT AATTCAATTT TTAGTTCAAT 2640

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CCATGCTAGT TTGATCCCAG CTCCCAAAAA ATAATTATTA TTCTTCGAAT AACTCTCAAA 2880

TACTATTGAC AAATCATTAA AAACGTCATT TTCAAAGCTA ATATAATATC TAAGACCAAG 2940

CAAAAATCTA TTGTCATCAT CATCAAATTT ATAATTTTTA GACCAAAGTA GGTCTTGTCT 3000

TAACGTAGAG TAAAAAAATA TTCCACTATA AATTTCAAAT AAAAAATCAA AAATAACAGy 3060

AATATCAAAA GTTTTATTTT CAATGCTAAA TGGGAATTCA ACCATTGCAA GTAAATCAAC 3120

TTCATTAAAA GATGCTTGAA AATAAAGCCA AGGGGATAAA TATTTGGGAG ATAAAAATTT 3180

ATCTAAGCTT TCTTTGTCTA ACATAGAACC CAAAGAGACA GAATAATTAG AAATAAAATA 3240

ATAAAACTCA AAAAACCATT CTGGTTCTAT TGTGGTCCTT TTTAAAACAA TATTTTCAAT 3300

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AATTCAAATT TATAAGGAGA ACATTTTGAA AAACCTTAAA TTAAATATTA TTAAACTTAA 4020

CGTTATTACA GCAATATTAA CTTCAATTTG CATATCATGT GCACCTTTTG GCAATGTTAA 4080

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ATAAAAATAA ATTTAATATT GTTATTTTGA ATTAAAACTA ATGTTTAGTA ATTCAAATAT 5040

ATAAGGAGAG CACATTTTGA AAAACCTAAA TTAAATATTA TTAAGCTTAA CTTTATTACA 5100

GCAATACTGA ATTCAATTTT C 5121

(2) INFORMATION FOR SEQ ID NO: 24:

(i) SEQUENCE CHARACTERISTICS:
(A) LENGTH: 5107 base pairs
(B) TYPE: nucleic acid
(C) STRANDEDNESS: double
(D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 24:

GCTGTGAAGG GTAATAATGA GAAAGAGAAG GCTGAGGGGG CTATTAAAGA AGTTAGCGAG 60

TTGTTGGATA AGCTGGTAAC ACTGTAAAGA CAGCTGAGGG GGCTTCAAGT GGTACTGATG 120

CAATTGGAGA AGTTGTGGAT AATGATGCTA AGGTTGCTGA TAAGGCGAGT GTGACGGGGA 180

TTGCTAAGGG GATAAAGGAG ATTGTTGAAG CTGCTAGGGG GAGTGAAAAG CTGAAAGTTG 240

CTGCTGCTAA AGAGGGCAAT GAAAAGGCAG GGAAGTTGTT TGGGAAGGCT GGTGCTAATG 300

CTCATGGGGA CAGTGAGGCT GCTAGCAAGG CGGCTGGTGC TGTTAGTGCT GTTAGTGGGG 360

AGCAGATATT AAGTGCGATT GTTAAGGCTG CGGATGCGGC TGAGCAGGAT GGAAAGAAGC 420

CTGCAGATGC TACAAATCCG ATTGCTGCTG CTATTGGGAA TAAAGATGAG GATGCGGATT 480

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AGGATAGTGT GAAGGGGATT GCTAAGGGGA TAAAGGAGAT TGTTGAAGCT GCTGGGGGGA 720

GTGAAAAGCT GAAAGCTGCT GCTGCTGAAG GGGAGAATAA TAAAAAGGCA GGGAAGTTGT 780

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TTAGTGCTGT TAGTGGGGAG CAGATATTAA GTGCGATTGT TAAGGCTGCT GGTGAGGCTG 900

AGCAGGATGG AGAGAAGCCT GAGGATGCTA AAAATCCGAT TGCTGCTGCT ATTGGGAAGG 960

GTAATGGGGA TGGTGCGGAG TTTGATCAGG ATGAGATGAA GAAGGATGAT CAGATTGCTG 1020

CTGCTATTGC TTTGAGGGGG ATGGCTAAGG ATGGAAAGTT TGCTGTGAAG GGTAATAATG 1080

AGAAAGAGAA GGCTGAGGGG GCTATTAAAG AAGTTAGCGA GTTGTTGGAT AAGCTGGTAA 1140

CAGCTGTAAA GACAGCTGAG GGGGCTTCAA GTGGTACTGA TGCAATTGGA GAAGTTGTGG 1200

ATAATGmTGC kAAGGyTGCT GATAAGGCGA GTGTGACGGG GATTGCTAAG GGGATAAAGG 1260

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AGGCTGGTGC TAATGCTCAT GGGGACAGTG AGGCTGCTAG CAAGGCGGCT GGTGCTGTTA 1980

GTGCTGTTAG TGGGGAACAG ATATTAAGTG CGATTGTTAA GGCTGCTGGT GAGGCTGCTG 2040

GTGATCAGGA GGGAAAGAAG CCTGAGGAGG CTAAAAATCC GATTGCTGCT GCTATTGGGG 2100

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GGAAGGCTGA GGGGGCTATT AAGGGAGCTG CTGCAATTGG AGAAGTTGTG GATAATGCTG 2760

GTGCTGCGAA GGCTGCTGAT AAGGATAGTG TGAAGGGGAT TGCTAAGGGG ATAAAGGAGA 2820

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GGATGGCTAA GGATGGAAAG TTTGCTGTGA AGAGTAATGA TGGTGAGAAA GGGAAGGCTG 3720

AGGGGGCTAT TAAGGAAGTT AGCGAGTTGT TGGATAAGCT GGTAAAAGCT GTAAAGACAG 3780

CTGAGGGGGC TTCAAGCGGT ACTGATGCAA TTGGAGAAGT TGTGGCTAAT GCTGGTGCTG 3840

CGAAGGCTGC TGATAAGGCG AGTGTGACGG GGATTGCTAA GGGGATAAAG GAGATTGTTG 3900

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AGGCAGGGAA GTTGTTTGGG AAGGCTGGTG CTGGTGCTGG TGCTAATGGG GACAGTGAGG 4020

CTGCTAGCAA GGCGGCTGGT GCTGTTAGTG CTGGTTAGTG TGGGGAGCAG ATATTAAGTG 4080

CGATTGTTAC GGCTGCTGGT GCGGCTGCTA GTGAGGCTGA TCAGGAGGGA AAGAAGCCTG 4140

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TGGCTAAGGA TGGAAAGTTT GCTGTGAAGA ATGATGATGA GAAAGGGAAG GCTGAGGGGG 4320

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GGAGTGAAAA GCTGAAAGTT GCTGCTGCTA AAGAGGGCAA TGAAAAGGCA GGGAAGTTGT 4560

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CGGCTGAGCA GGAkGGAAAG AAGCCTGCAG AkGCTAmAAA TCCGATTGCT GCTGCTATTG 4740

GGAAtAAAGA TGAGGATGCG GATTTTGGTG ATGGGATGAA GAAGGATGAT CAGATTGCTG 4800

CTGCTATTGC TTTGAGGGGG ATGGCTAAGG ATGGAAAGTT TGCTGTGAAG GGTAATAATG 4860

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ATAATAAAGA GGCAGGGAAG TTGTTTGGGA AAGTTGATGA TGCTCATGCT GGGGACAGTG 5100

AGCTGCT 5107

(2) INFORMATION FOR SEQ ID NO: 25:

(i) SEQUENCE CHARACTERISTICS:
(A) LENGTH: 5068 base pairs
(B) TYPE: nucleic acid
(C) STRANDEDNESS: double
(D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 25:

CACATTATTA AATCTAGAAT TGAATAATTA TTCTCAAAAA AAACTATTAA AATTTTACAA 60

CGAAATTCTT AAAAAAGATA ATAAAAATTC TTGCGATCTA CCAACAATGA ATAAATATCT 120

TGATATATTA GAAAAAACAA AAACCATAGT AAAACTATCT TTTAAAAACC AGTCCAAATA 180

TATGATTTAT TATAAAATTA ATTACCCCCT TAAAGTGTTT CGTTCAACAA TACAAGACTA 240

CTATCAAACA ATArCAGATA AACTAAAACT ACGGTTAGAA CTAAACTATC CTACTACTAT 300

TTAATCGTAA AAAATATTTC TTTGCAAATT AAGCAATTTA GAAATATAAA TGTAAAGACA 360

TATATTTTTA TTTGATAAAT AATAAAAATT ACTGGGGCAC TATTTGGAAA AATTTTTAAA 420

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ACGCCGTGCG TTTACTTTTA ACATAGATAG ATATTGCAAT ACTAGTAAAG ATATTGAAAT 600

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TAATAAATAT AAAGTAGAAA AAAATATATT CAAACTTTAC TACATAATCA ATTATCCTTT 720

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TTGCTATATA GTTTACTTTA TGAAAAAAAT CTATCTATTT TATGCAAAGA AGATTAGTGC 4560

CACCCTGATT TATATGATAA TTGCCTTCCA AATTAACAAC ATATCGAATC ATTAATACTT 4620

TTCCCCAAAC ATTCATTTTT AAATACCCTT AATTGATAAG CTCTCTATTG TCAATGTCCA 4680

AGTGCTGCCC TGCTACTAAA ATGCAATATT TATTATAAGC AACTTTCTCT ACTACTGTAT 4740

TTTTATTATA TATTAAACAA AAAATCAAAA ACCTAATAAT ATATTATTTT AGGTTCCCCC 4800

AAGCATTCTT GATAACAAAG TTATTTGATA TTGGTTTAAA TTCTTGTAAA TTATTTGTAC 4860

ATACCATTTA CTTTATTATA AAAATAAAAT TTCCCTTCAA TATATTTCCC CCACTTATCT 4920

AATTTAATTC TCTGTGTTTT TAAAAGCATT TTTGTTTTTC ATGTTTTTTT GATTTTCAAT 4980

CATAAATAAC AATAAACTTA TTTATAATAT TGAGAATATT ATCTAATAAA ATATTAAAGA 5040

TGTAAAAATT AGTTACAAAA AATTGCTG 5068

(2) INFORMATION FOR SEQ ID NO: 26:

(i) SEQUENCE CHARACTERISTICS:
(A) LENGTH: 4663 base pairs
(B) TYPE: nucleic acid
(C) STRANDEDNESS: double
(D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 26:

TTCTTTATTT TGGCATTCAA CTTTACATTT ACCTTTTTTT ACAGTTTCAA TAGGTTCTGG 60

TGCATTTTCC ATGTTAAATC CTTATACGGC CTTTATGCTA AATTCTTCTG TGGTTAAAGA 120

ATTTTTTTCA TTTTTTATTA TTTCTAATAA TTCAAGATGA TATATACCAA ATACTTTATT 180

GTATTCTATT TTCTTTTGGT TATTCAAATA TTTTTTGATA ATTGGCTTTA GAATTTCAAT 240

ATTTGTTTCT TTTTTTAGTT GTTCAATAAG GATATTGAAA ATATTTTCCT TTAAATTTTG 300

ATAATTTTGT TGAGAATTTT gCTTTTyycT TTCAATCGmC TTTTCTAATT TACGTTTTAT 360

GTTGTTTAAA TCGyTATATT TATGmyTTTC AATAATAAAr TGGGGCTTAA ATTTGTAAyT 420

TTCGTATACT TTTTGTAAAT TTrTCTTTAA TTGTTCGGGr TTATATCCAT TTTGTTyGAA 480

TTCTTTCTGA GTGTTATTTA rAATTyTyTT TAGTTTCTCT TGCTTTTCTT TAAAGCAAGA 540

TTTATTTAAA TTTGTGTTTT TTAGCAATTT ATTTTCAACT CTTTTTATGT TTTTGATTAT 600

TTTAATCATA GTATCTTTAT CAACATTTAA TGTTAATAAA AAyGGAAGAA TTTCTTTACA 660

TAAsAAGTTr CTTTTrTTGA ArTACTTTAT TACTTGATAT TTTTCTATTT CTTTAATCTT 720

TCTTTCTTCT TTTATATTAT TTTTATTACT TAAACACTCC ACTGAATTTA CACTACyATT 780

TTTAGAAAyA TyGTCTTTAA AATGGTTATT AACTCTAGAT TTAAATCTAG AGTTTTTTCG 840

yTCTTyAAAG TACTTGTTGA TTTTCTGGTA ACATTCTTTT TTAGGATACT TTAGTTTATA 900

rTAAATTTCA GTyCCACAAT TTACACCCAT rTGTTGGTAG TAATTAGTTG TrACTTTTAr 960

TACTTTTTCT AATTTrTAAA GATAATTTTG CATTGTTCTT AGwGTAGTGG GAGCTAGmCC 1020

ATTTCTTTTT AGATTTTCrT TAAAGCAGTA GAGTATGTTT TGTTGTGTGT ATTTTTTATC 1080

TTTTTTGTTT AGAAAATCTA GCGTTGAAGT AAGAGATATT AATTTGTGTT GGTGTTTGTT 1140

GTGGCAAGTT GGAATTTTTG TTTTATTTGG GAAATTTTGC ATTTTTTTTC TCCGTTTACA 1200

TGTATTAGTA ATTACTAATA TAGTGCAAAA TTTTGATTTA AAAGTAAATA CTTTTCTAAA 1260

AAAATATTAA ATTTTAATTA TTAATTTTGT TAAATTAATA TAAAAATTTG TAATTTAGTA 1320

AAAATATAAA TTGATTTTAA TTCtAAGTTG ACTTATACTA CAAATAGCGT AGTAAAATGT 1380

TCGCATTTTT TCTACCGTTT ATATGAATTT ACAAGAAATA AAGATATGGG CTTAACTAAG 1440

TTCTTTAACA AGAGAATTTA GCTAAGCCCG CACTTTTTGT AAAAATTTTT TGTAAAGAAG 1500

TTGGCAAAAA TAGTTTTTGC TATATACTTA TATTTATTAA TACATATAAA CGGAGGAAAA 1560

AGATGGAAAA TCTTTAAAAC AATAATAATC CACAAGAAAA TATTCAAGCA GAAATTAAAT 1620

TCTTAAATGA TATGAAAACC CTAATGAAGA ACTTGCCACG TATTGACAAA AGTCTTAAAG 1680

GGTACGGATA TAAGTATCAG GATTTCAATG ACATATTAGA AGAAATTGAA AACGTTATTG 1740

AAAAGCATAA TTtGGATCTt AtGTTTAGGC AATTTCCAAC TTTTACACAT GATCCATATG 1800

GTATAGTTCA TGTTATTAGG ACCACATTCT ACAGCACAAG CAGTGGGTAT AGAGAGTCAT 1860

TTGATACCCC AATTCTTACA GAAAATTTGC AATGGAACAG TGAAAATGGT TCTAAAAATG 1920

TAAATACATT GCCACAATTT GTTGGTTCAG CTATTACTTA TTTCAAAAGA TACGCTTTAG 1980

TTGCATGTCT TAATATAAAA AGCGAAATGG ATACTGATGC AGCACCTATT TACAATAATT 2040

ATGAAAACAG AAATTCTATG CCTAACAAAC AAGTTAGTGT TAATCAAAAG TAAGAACAAA 2100

AAAGAGAACA AAAACAAGAG ATTAATCAAA TTCAGAAAAA TAAyAyTATT CAAAACCAGA 2160

AAAGAGATAT TAAGCAAGAA TAAAAAAAAG ATAGGTTTTA TTATTACAGT GTTTTTAAAG 2220

AAGCGTTGTC TAATATAAAA GATTGGGTAA ATAGCCCTAC AACAAAAGAT AATATAAACT 2280

CAATTATTCA AAAAATAAGC TTTATTCAGA ATATAGACCC CAATAATGTT GATGATATCA 2340

AGAAAATTGA ATCTGATTTA ATCTCGTATT TTGAGAAAAA TAGTGATTTT AAAAGTATAA 2400

ACTATTGGGC GGAGATTATA AAAAACTATT TCAAGAAAAA TAATArATTA AAGGATTTAC 2460

AAGATTTTGA AAAGTTTGTG gTgTTTAAGA GGACTGCTTA TGGTCCTAGC CCATTAATAT 2520

TCTTTAGTGT CTTAAAAGAA TATGAACGGT TTGATGAGAT ATTTGCAGCA TAGCAAGATT 2580

CTTACATGGT AAAGCCCCCC TATTTGGGGG CTGCTATATA TTATGAATTT TTGCACGTAC 2640

TACTTGCAGT ATTTTCAAAG CCGTCTATGC CCCCTTTAAG GGCCTCCTGA ACGACCTGCT 2700

TGAAGGTATT TTTATTTTCA ACCTTATCTC CAGTACAACT GTCAAGTTCA CTCTTTATAT 2760

GATCAAGTGC AGATTTTATT TTGCTTTCAT CATATCCTAA AAATTTATTA AATTCTCCAT 2820

CATTGCCCAG agCTtCTTTT AACCAGTCAA GrTGTGTTTT TTGrTywTCA kwTAGCTTTT 2880

CTCTAAGmAG kTCTTCTTTA GATTTwGGTT TTTCTTGTGT TGCTTCTTTT TGGGTTAAAT 2940

CACGTTTTTG TCTACTTTTT GTTTGGCTAk TATTAGTATC ATTAGAATTA CAGCyGTTTA 3000

GCATTAGTAA AAATAAACAA AATAATATGT TGATAATTTT CATTrTTATT CCTTTTTTTA 3060

TTATTAATAT TCACTTAATC AATTATTAAT ACTAAATATT GGATAAACAA TTATTATTTG 3120

AATTGATATG TTTTAAGTGA GGTAGTAGCT ATTTAGAAAT GAAAGCAAAT ATTAGCCCTG 3180

CTATCATTGT GATAGACATT GCTCCCATAA TTCCCAATAC CCATTTAAGC ATTTCTGAAA 3240

GAGACATTAA ATTCTTTTCT ACATTGTCTA TTTTAGCAGT AAGTTCATTT TTAACACTAT 3300

CGATCTTAGC ATTAAGTTCG CTTTTAACAC TATCTATTTT GACATTTAAA TTCTTTTCTA 3360

CAGTATCTAT CTTAGTATCT AAACCATCTA TTTTTAGATT TAAATTCTTT TCTACAGTAT 3420

CTATCTTAGT ATCTAAACCA TCTATTTTTA GATTTAAATT CTTTTCTACA TTGTCTATTT 3480

TAGTATTAAG TTCGCTTTTG ACACTATCTA TTTTAGAAAT AAGATTATCA AATTTTATAT 3540

CAAATTGTTT TTCTAAATTT TCTAAATCTC TATATGTTAG CTCATTGTGA TAATATCTTT 3600

TAGATAAATC TTGTGCTATT AATTGTTCCA TGCCCAGTCT AATAAATTCT TTATATATTT 3660

GTTCTTGAGT TACACTTGCA ATATTTGTTG ACACTGTTTC CATAAAATTT TCCCTTATGG 3720

TCATATTATA TACTATTTTA GATTAATTGG CTTTAGAGAT TTTTATATGT AAAGTAGAAT 3780

TTCTTGCAAG AAAAACCTTT TTGTAATTTA CATTTTTAAC TTCAGATATC AGTTTTAAAT 3840

TTTTTACTGT AGATTTTTTA CAAAAACAGT ATTGCAAAAA CTCTTAGATT ACTTTTTCTT 3900

TTCTTTGTAT ACTACAATAA CTCCAAAACC CACTAAATGG TTTAGTGATT TAACCTCAAG 3960

AATATTTTCG GCACCTATTT GGTTAATAAA ATTTTCTAAC CCTATCCCTA TAATTTCGAA 4020

TAGAGTTTTG TTTTTATCTT CTTTTTTTAT AGGAAAGTTA ATGTTATGCT TATGATCATC 4080

ACCGCCTTGA TCTAAAGCTA TTAAAGTTTT AACTTTTATA ATTTCATCTT TTTTAATTTC 4140

ATATGAAATT AAATTACCAA TACTGATAAT AAACATAAAT AACATTAATA AATTAATTTT 4200

TTGCACATTG TGTTCCTTAA TAAATAGAAT ATTAACAATA TTATATCTTT ATTAAgATTT 4260

GCCCTAAAAT ATAAAATTTT ATTAAAATAT AGCAGTAATA AACGACTTTA AGAATATAAA 4320

TGGGAATTTC TTGCAAGAAA AAcCTTTTTG TAATTTACAT TTTTAATTGA GAATATTTAT 4380

TATAGACTTT TTCCGCTATT GGTTTTGTTT TTTTAATGTA CTCTAAATAT ATGTTGATAT 4440

TATGTTTTAC AGCAGTTATG GaGTGTTCGT CTTTTArTGT TGATAAATCT GGaTAAGGAT 4500

ATTCTGGaTA ATTTGGaTCA TTAACTTTAA CTTTTGTTTT aGCTAAAAAT GTTACTAGGT 4560

ACATAACATA CTCTGAAAGT TGTGTTTCAT ATTTAGCTAA AGATTTTAGT GTTTGAATAA 4620

TTGGAGGTTT TGGCTCTTCT GGTAGGTTAG CAATAGTGGT GCA 4663

(2) INFORMATION FOR SEQ ID NO: 27:

(i) SEQUENCE CHARACTERISTICS:
(A) LENGTH: 4312 base pairs
(B) TYPE: nucleic acid
(C) STRANDEDNESS: double
(D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 27:

CAACTTCAGT TTATATTATC AAACGACAAA TAAAACATTA AGTACAATAA CCCAATATTT 60

CAAAAAGGAT GCACATTTTA ATAAAAAACT AAAGCTGTTC AACTACAATT GTTGCACTTG 120

AAATTTTTTA TACTAAAATA TAATACAAAT AATTATATTA ACAAATATCG ATTTTTATAA 180

AAAATAAGTA AAAGTAGTTT AGTTTACCTG AGTATTTAAA TACTTTTAAT TGAGGATGTT 240

TTATTTTAAA AAGGAGTGTA AAACTATGTC AAAAACTGTT GACGAAGTAT ATTGCTATTC 300

TTGTGGCAAG ATTTAAAAAA GATGCTGAGA TTTGTATTTC TTGCGGGGTC AGAAATAAAC 360

AAACCGAAAA CTACAATAAA CTTATAGTAT TTTTACTATG CTTACTTTTT GGTTATTTAG 420

GAGTTCACAG ATTTTATGTA GGTAAAATAG AAACTGGGCT ATTATAATCA TACCTCTTCA 480

CTTTTAAATA TTGTTCATAA GCAGTGGGTC TAGGCATATT ACGATTATAC TCATGGCTCC 540

CCTCGCCAGA ATACTTAATA TCTAGAGAAT ATAACTCCTC TATACATGAA TACAGCCAAC 600

GAACTATACG CATAAATTTA TTATTGTTCT GCTCATTTTT TACCATAATA AACAACTTAA 660

TCAGGTCTTT AGATTGCTCA GACCCTAAAT CTAGAAAGAA TTTATTAAAA TACTCCTCAT 720

TATAGTCACT CACTGGGmGT ACTACAyCAC GATAATTAGG ATCTTTTAAA CTAATTAAAG 780

TCTTTTTTAT CTCAACCAAT AAACGATGTT CAGACGCATA TAATTTTAAA CTAGAAAATA 840

TGCCTTCAGT AACAATCTCA TCGGAACCCA TATCACTACT TTCCGAATTC TCAAAAAAAG 900

TTTGAACAGC TTGTTCCTGA TTATCAGGCA AATAACAACC CAATAAAGCC AATGyTATTA 960

TACAAACGCC AACCAAAATC CTCATATATG CTCCTTACCT ACCTTACTAC AAAAATAATC 1020

TGTTGTAAAT ATACAAGTTT ATCTAAGCAC TATTTTATTA AACATTTAAA GTCCTACCAA 1080

GATAGATTTT TCTATAACAA GTAAACATAT ACCAATTTTA CCTTTTCTCA AGAATTATTA 1140

AATACTAAAA TATTAATTTT AGCTTTATTA TTATCTAGTG AACTGCTATT TCTAAATCAA 1200

AGATTATAGA AATAGCAGTT CACACAAAAT GTGCTTAAAA AATTAAAAAA TAATTTTAAC 1260

AAGACTAATA AAAAATTTAC TAAAGAATCC TTTTAATTAC ATTTAAACTT GTTTGTTGTT 1320

ATTCTAACAA GATCGATTAA AACTCCAACA TATAAAAATC CAAATGTAAA TAGAGAAATA 1380

GAAAGAAATA CAATTAATGG AAAAATTTTA TGAAAAAGAA CATATATATT TTGAATATAT 1440

TTTTATATAT ACCATTATTT TATTCGTGTT TTTTGACTCC ACCAAAATCT TCAAAAATCA 1500

ACAGTATCAA AACCGAGGTT TTGGATTTTA AGATAATTGA AGAGGGAAAT ATTATAAAAT 1560

ATGATAAAAA GCCCATTGAA GAGCGTAATG AAAATACTTG TCTTTCTTTT AAAGAACCCG 1620

AATTAAATGA AATAAAAGAG GGGGACGTGC TTGAATTACT TGCAGGTGGT TATGTTACAT 1680

GGGCAAAATC TGGTGACTTA AGGGTTTTAA AAGATAAAAA TAACAATTTA ATTGAAGATC 1740

TTAGAGAACT TAGGTACTCT TATATTTTTT CACCCATTCG ATTCAAAACT TTTTTTAGTT 1800

ATAATTATAG CATTAATGAC AATAACTATA AAATACTCGG CAAAAAAGCA CCTATAGTTA 1860

AGATAATAGC ATTTGAATCA ACTAAAGAGT TTGAAAAAAA ATACGAAATA AATAGTTTAA 1920

AACTAAATTC TGAAGAATCT AATATTGATT TTGAACAAAA TAGAACTGGT TTAGCCAAAA 1980

TTAATTTAAA AGAAACTTCA AAAGAACCTA ATTACATTTA TTCATATAAT TTTGGAGTTT 2040

TTGACAATTC TTTAGCAGAT TATTTTAAGT TATTTTATAA AAAAAATAAC TGTAACTATA 2100

TGCCTGCATA TCTTACTATA AAAGATAAAG AAACCGATAA ATATAAAACC TACGAAATTA 2160

TATTAAATCT AAAGCTATTT AACGATACCA TTAAATTATT AATTAATAAG TATTCAAATT 2220

TATCAAAAGA AAAATTAAAA CTTTTTACTG ATGAATGATA AAAATTGAAT AAGAAAAGCA 2280

AAGACAAAAA TTTATCAATT AATGTAAATA AATACAATTC AAAATTGATA AAGTTGTGTG 2340

ACATATTCGG TATCTCAACT TGTCGATTTA AAAGTATTGA TAGTGAACTG CTATTTCTAA 2400

ACAAAGATTA TAGAAATGGC TTCTTGTATC TTCCCATGAT AGCTTTTTAG GCCTTTTGTG 2460

CAAGAACCTT TAGTTAAAGA AACTATTACT AAACTAGTAA TAGCAGTTAA ACTATAAAGA 2520

ACCTATTTTT CAAGAATAAA TATTTTTGAT ATAGATTCAG CAATACATAA CTCTAAAATA 2580

CTATTTAAAA TCAAAATAAA TTTATTTATC CTCAAATTCA TCTCATTCTA ATTTTTATTT 2640

AAATTCTTAT TCTATGTTAT TTAGTTTTAT GATATCTATC TTAACATCTA GCTCATAATC 2700

TTGATTGCTA CTATATATGT GATATAATGA TAAAATATTC TAATAATATT CTATTTTAGA 2760

TAGAGGTAAT ATAATGAATT TAATAATTAA AGTGATGTTG ATATCCAGTT TATTTTCTAG 2820

CTTTATCTCT TGTAAATTAT ATGAAAAGCT TACAAATAAA TCGCAACAAG CTTTAGCTAA 2880

AGCTTTTGTC TATGATAAAG ATATAGCTGA TAATAAAAGT ACAAATTCTA CTTCTAAACT 2940

AGATAATAGT TCTCTAGATT CTATAAAAGA CAACAACAGA AGTGGTCGCA CATCTAGAGC 3000

TTTAGATGAT GCTGAAGAAA TTGGGGTAAA AGAAAGTAAT CAAAACAGAA ATGATCAACA 3060

ACAAAATAAT GAAAGTAAAG TAAAAGAAAG TGAAAAAAAC AATAGTTCAG GTATACAAGC 3120

AGATGATAGT GTTTTAGACA CAGCTCATTC CGATGCTAGT GAAGTAGAAA ACAAGAAACA 3180

TGATACTAGC AGACAACCTC AACTACTTAA TAAGGACTCT AGTGAAGCTA GAGAAGCTAG 3240

TAAAATTATA CAAAAAGCTT CTACCTCTTT AGAAGAAGCT GAGAAAGTAA ATGCAGCTTT 3300

AAAGGAAACA AGATCAAAAC TTGATAAGAT AAAAAGATTA GCTGATAGCG CTAAATCTTA 3360

TTTAAATAAT GCTAGAAAAA ATTCTAGAAC TAATGGTTCT ATACTAGAAA TATTGCCTAA 3420

CCTTGATAAA GCAATTGAAA AGGCTATTAG TAGTTATGCC TCTCTTAATG TTTGCTATAC 3480

TGATGCAATT GCTGCTTTAG CAAAAGCTAA GAATGATTTT GAGCATGCAA AAAGAAAAGC 3540

AAATGATGCT TTAGAAGAAG CTTTAAAAGA TATACCTCAT TTTAGGGGGT ACAATTATCT 3600

TTACCATTAC CGGATAAATA ATGCTAATGA TGCAATGGAG AGTGCTAAAA GTTTGCTAGA 3660

GGTTGCTAAG AATAAACAAA AAGAACTTAA TGAAAATATG ACTAAGACAA ATAAGGACTT 3720

TCAAGAGTTA AATGATATAT ATAAAAAATT GCAAGATATG GACTCTAGAT AAGTAAAAGT 3780

AAAATATTAA AGATAAGAAG CCAGACAATA CTTTATGAGG TTTAGCTTCT TTATTTATAA 3840

TATTCTTTTT CTAAACAACA CTTTATTTTC TCTTAACTTT ATAGTTTGAC TTAAAAAATC 3900

ATTATTTTTT AAAATATTAC ATGAATTAAC TTTAATATCT TTATTTTTAT ATTATAATTA 3960

TTATTAATAT AGATATTATT TGCCATATAA GTACATAACA AAGTTTTATT AAAAAGGAAA 4020

TATAAATATT ATGCGATTAT GTTTAATAAA AATTTTTATT ATACCTAATT TAGTATTTAG 4080

TTCTCTTTTT TTATTTGAAA GTTGTTCTGG TTTTCTATCT AAAAAATCTA TAGAACAGTT 4140

TGCATTAGCA TTAAAAGATC ATCAAGAAAA TAAAAATACT ACTAATACTT CgTAGaTAAA 4200

AATAGTAAGG AAATTGAATC GCCTAAAGAC GTTACATCAT CAAATAAAAA AACTTATGAT 4260

CCAATCTTAC AAGTAGGTTC TAAATCCCCC AAAAAAAAnC CCCnAAAATT TT 4312

(2) INFORMATION FOR SEQ ID NO: 28:

(i) SEQUENCE CHARACTERISTICS:
(A) LENGTH: 4305 base pairs
(B) TYPE: nucleic acid
(C) STRANDEDNESS: double
(D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 28:

nAATTCGAGC TCGGTCCCCA AAAGTAAAAG CAAGCCTTGA TAAGATAAAA AGCTTACTTA 60

GTACAGCTAA ATCTTATTTA GAGCAGACTA GAAGAGGTGT GGGATCTAGT AAgCTAACTT 120

AGCCTTATTG CCTAGCCtAG AAGAaGCTAT TGCTAAGGTT AAGAGTAATC ATGCTTCTGC 180

TGATACTCAT TGCAATGATG CTATTGCTGC TTTAAAAAaGA GCTAAGAACG ATTTTGAGTA 240

TGCACAAAGA AAAGCAGATC GGGCTTTAGA AGAGGCATTA AGTAATAGCA ATGCTTCAAG 300

GCATGAGAGC TACTACTATG CTGGCTACCA CCAATTTATG GCTGATGCTA AAGCTTCAAT 360

GAGTAGTACT AAAAGTTTGC TAGAGGTTGC TAAGAATAAG CAAAAAGAAC TTAATGAAAA 420

TATGACTAAG ACAAATAAAG ACTTTCAAGA GTTAAATGAT ATATATAAAA AATTGCAAGA 480

TATGGACTCT AGATAAGTAA AAGTAAAATA TTAAAGACAA GgCCAGACAA TACTTTATGA 540

GGTTTGGCTT CTTTGTTTAT AATACTCTTT TTCTAAACAA CACTTTATTT TCTCTTAACT 600

TTATAGTTTG ACTTAAAAAG TCATTATTTT TTAAAtAATT ACATGAATTA CCTTTAATAT 660

CTTTATTTTT ATATTATAAT TATTATTAAT ATAGATATTA TTTGCyATAT AAGTACATAA 720

CAAAGTTTTA TTAAAAAGGA AATATAAATA TTATGCGATT ATGTTTAATA AAAATTTTTA 780

TTATACCTAA TTTAGTATTT AGTTCTCTTT TTTTATTTGA AAGTTGTTCT GGTTTTCTAT 840

CTAAAAAATC TATAGAACAG TTTGCATTAG CATTAAAAGA TCATCAAGAA AATAAAAATA 900

CTACTAATAC TTCAGTAGAT AAAAATAGTA AGGAAATTGA ATCTCCTAAA GACGTTACAT 960

CATCAAATAA AAAAACTTAT GATCCAATCT TACAAGTAGG TTCTAATCAA CATATGTCAG 1020

ATGATCCTGG TGCTAATAAT AAAGAATCCC TACCAAATTC AAGTCCAGCA ATAATACAAA 1080

ATGACTCGCA TGCTCAAAAT AATGTAAAGA TGGAAGAAAA TAAATCAGCT ACTCCACAAC 1140

ATGATCCAAT TGAACAAAGT AATTTTAAAA ATAGCCTTAC TACAACAAGT AAAACTCCTG 1200

CTATTCCTTC AGAAGAAGAA ATTAAAGCTA ACTTAGATGA ATTTGCACAA GAAGAGTATG 1260

AGCAAACATC TCTTTCAGAA ATTAAAAATG CCACGCAAAT TGTTAATCAT GCTAATCCTG 1320

AAAACAAATT AAACAATACA CTCCTTGAGT TTGAAAAAGA TTATGAAACT TTATCAAACT 1380

TGTTATTCTC TAATTTAGAC GCATCTCCTT TGAATAGAAA AATAAAGACT ATTATGCCTA 1440

AATTACAAGA AATGCGTTCT TTTATGGAGC AAGCAACTAA TTCTTGGGTA TCTGCTAAAG 1500

GCATGCTAGA TGAGGCTAAG GATAAACTAG CAGAATCTAT TTATAAAAGA CTATACAATG 1560

GCAATTCATA CCGGTTCGGT GGCAGTTTTA ACGGACGTGA TATGCAACAT GCAAAAAATT 1620

TAGCATACAG AGCTATAGAC TTTGCTTCTG CATGCATTGA ATATACACAA AAAGCTATTG 1680

ATTATCTTCA ACAGGGAAAT TCTTGCAAAA AAGAAATAGA AAATATATTC AAGCTTTAAA 1740

ACTTCCAGTG TAGGCTTTAG TTTCTTTAAT ATCTCTACTC ATATATAATC CCATCTTTAC 1800

TAGAAAAGCT TATATATCGG CTTACCTAAA TTAACTATTT CACTTCCCTC TTGCCTTTAC 1860

AAGCACTACT CTACTTCTTT CAAATTTATA GTTTGGTTTG CATTTAGCAT TCACTATTTT 1920

TCTATTTTTA TAAATGTGAT ATATTTATTT TTTAAGAATA AAGCATAAAT ATCATCAGAT 1980

TCTAAGAAGA GGTACTAAGA TAGATGAATT TAATTGCTAA ATTATTTATT TTATCCACTT 2040

TAGTTTCAAT TCCAAATATC CTCTCTTGTA ACCTATATGA TAATCTTGCA GACAACGCTG 2100

AGCAGGTTAC AGACATACTA GACAACAACA AGTCTTTTAA TACTTTAGGA AGCAGCAATG 2160

AGAGTAGAAG TCGCAGGCCT AGAAGTACAA ATAATGCTTA TATGAAACAA AACATAGACA 2220

AAAATCATTT AGTTGTTGCA GATATGCAAA ATGATAATAG TAGCAGCAGT CTTCCCCAAC 2280

AAGTTAATAG TGAATCCAGT AAAGCTAATG AAGATAGTAA TATTATGAAG GAAATTGAAT 2340

CTTCTACAGA AGAGTGCGCT AGACTAAGAA AAGATTTAGA AACTATAAAA CAAATACTTG 2400

ATAATATAGA AAGCTTGCTT AATACAGCTA ATTCTTATTT AGAGAACGCT AGAAAAGCAC 2460

CTAAATCTAA TCAAGATAAT CAAACCTTAT TGCTTAGCCT GCACCAAGCT ATTGCTAAGG 2520

TTAAGAGTAG TCATACTTCT TTTATCATTT GTTATAATGA TGCATTTAAT TCCCTGGGAA 2580

TAGCTGATAC TGCCTTTAAA GATGCAAAGA GAAAGGCAGT TGAGGCATAA AATGCTTCAA 2640

AGGAAAATTA TGAATGGTAT AACGGTCATT ATCATTCTTT TATAAATGAC GCTAAAGATG 2700

CAATGGAGAG GGCTAAAAGG ATGCTAGATA ACGCTAAGCA TAAACAAGAA TATCTTAATT 2760

CTAATATGTA TCAGGCAAAT GCAGACTTTG AAGAGCTAAA TAAAGCATAT GAAGCTGCTT 2820

ATTAAATAAT ACTAATCTTA GATAGCTCAG CTTTAAAGAA AAAAGCTCTA TGCTATAAGC 2880

TTAACATATA ACTCATATTC TTGATTACTA ATAATAAATG TAATATAATG ATATTGTATT 2940

ACAATAATAC TCTATTTTTG CATAAAGTTA GCACAATGAA TTTAATGATT AAAGTATTAA 3000

TATTCAGTTT ATTTTTAAGC TTTATCTCTT GCAAGCTATA TGAAGCTGTA GATAAATCTC 3060

TTATAAAAGA CAACAAAAGA AGTGGTCGTA AAGCTAGAAG TATTAGTTAT AAGGAAGTAA 3120

ATAATCAAGA ACAAAATAAT GAAAAGAACC TAAAAGAAGC AAAAGAAAGT AAAAAAAACA 3180

ATAATTTAGG TATACAAAAA GATGGTATTG TAAACACAAA CCCTTCCGTT GCTAGCGATG 3240

CTAGTGAAAA ACATACTAAT AGACAACCTC AACAAGTTAA TAATAACTCT AGGGAAACTA 3300

GTGAAGCTAG AAACATTATA CAAGAAATTT ATACCTCTTT AGAAGAAGTT AATAAAATAA 3360

CTACAGATTT AGAAACAATA AAGTCAAGAC TTAATAATAT AAAAAGTAAA GTTGACAATG 3420

CTAGTTCTTT TTTAAATAAT GCTAGAAAAT CTAATAAAGC TAACCCAACC TTATTGCCTA 3480

AACTTGATCA AGCTATTCGC AAAGTTAGTA GCAGCCATGC TTATGCTAAC TCTAATTATT 3540

CAGATGCAGT AAGTGCCTTA AAAAGTTCTA AGCACGATTT TGAGTATGCA AATAGAAAAG 3600

CAGAAGATGC TTTACAAGAA GCGTTAAATA ATAGCAATAC TCAAGGTTAC CAATATGCTC 3660

GATACCACTA TTATATGAAT GATGCTAAAG AAGCAATGGG CAGGGCTAAA GTTAGCCTTA 3720

AGACTGCTAA GCAGAAACAA GAAAAACTTA AAGACAAGAT GGATCAAGCA AATAAAGAGT 3780

TTGAAGAGTT AAATAAAGCA CATGAAGCTG CTTTAAGTAG TAGAGAATCT TAGCTGGTGT 3840

AGCTTCAAAG AAGACATGAC ACTTACTTAT AAATAAGGAA GCTTTTGGAT TTTAACAAAA 3900

ATAGTCTGGC TTTTTTGCAC ATATAAAACA ACTCCATTAT TTCTAAGATA AATATTTTAA 3960

GCTCCCTGGT AAAGTAATTC ATTTATCCTA GATTTACTCT TCCACTTCTA TACGTCCCGT 4020

CCTGCTTAAT CATTAATTTT TAAAATTAAA TGTTTCTTTC TAGTTACGCA CTATATTGTT 4080

ACTATAACAA AAATTGAATC TTAAAAATTA ACATATTACT TTAAAAAAGT ATACTTATAG 4140

GAGATGCTTA TAAAGCTTAA CAAACTTATT TTTACCAATA TATATATCTA ATATCTCTTA 4200

TACTTAGTTG CTCAATATCT AAAGATTTAA GTGATAAGCT CTCTTCACTA AAATCTAATG 4260

ATTTTTTTAA TTCAGACACT TTAGTTTATG ATTCTAGCAA TTATG 4305

(2) INFORMATION FOR SEQ ID NO: 29:

(i) SEQUENCE CHARACTERISTICS:
(A) LENGTH: 4293 base pairs
(B) TYPE: nucleic acid
(C) STRANDEDNESS: double
(D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 29:

CGATAGCTTT AATAAGcAAG CTCAATTACA ACAwTTAAaC CCAGGTGTTT CACAAAAACA 60

AGAwATAGAT AAAAAAACAT AGAAAAAAGA AGCCTAGACA ATTCTTCACA AGAAAAAGAA 120

CTCACAAACC CTGCTTATTC AACACAAGAA CATACAAAAA GTGCTACAAA CTTAGACTCA 180

AAAAAAGATG CTCTTATTAA AGAAACTCTT GAAGCTATAA AGGAAAAAAT TAAAGAAGAA 240

AAGAAAAGCT ATTCTAGAAG AGCAGCAAAG ACAGAAACAA CAAGAGCTTG ATAAGATTAA 300

AGCACAATAT GAGGAAGAGA AGAGAAAGAG AGAAGAAGAG AGGAGAAGAG AGGAGAAAGA 360

GAAAAGAAAG CAAGACTTCA AAAATTCATG CAAACTACTT CTGACTTAAC TAATCTTGTT 420

AAGATGGCTG GGCTTGAGGC TTATAGCATT TCCCATAAAT TAAAAGATCT TGAAAAAGGT 480

ATTGAAAATT ATGAAGACAA CAATAATTCT ACTAAAGACA CACTAAACCA ATCTCTTAAA 540

GATGTTATTT aTGAGATTAC AAAGCTTAGT AGTCTTATAG AAGCAAAAGA TAAGATTGAT 600

CAGCGTAAGA AATTGGGTTA TCAGACAGAA CAAGAGTTTG ATGCTAAATT TATAAACTTA 660

AAGAACATCA AAGATAAGCT AAAGACTTTA TGTGGTAAGG CTAAAGGCCA TCTTGGTAGC 720

AATCTTTCTA GCGTTACTAT TGATGGGATT ACTAAAGAGA AGGTAGCTCm AGCTTATCTT 780

ATCATTAAAC TAATACACAA AACATTAATT TATATGAATG ATGATAGTAA AGGTAGCCTT 840

GCTACTATAC TTAATGACTT AGAAAAGGAT GCCAAATCAA TATAACTAGC ACAACAATAT 900

CTTCTTATTT TAAAAAAGCC TAAGTACTTA TATCTTAGGC TTTTTTAAAA ATTATCTTGC 960

CTCTTAGACC ATTCTTGATA ATAATAATAC TGTTAAGAAT AAATTAATGC TAAAATGGAT 1020

AAGTACACTT ACACTAATTA CTATTTTTGC AGTAAATATA AAAACATAGA ATAAAAATTC 1080

TTATAACTAA AAGAGTATAA TCTTCTTCAA GAGAAATATA TTATCAAGAA ATTAAATAGA 1140

CCTTAAAATT ATTTTTTCTA TATATTCAAA ATTACTTTAA CTATAATATT ACCTATAGAG 1200

AAAATCGTAT GGTAGTGGAC GAATGGTTTT TCCTAAAAAT AAAGAAAAAT AGTAAAAATG 1260

GAAGAAAAGA AAACAAAAAA AAGATATAAT TCTGTATAAA AAAATTCTTT AAGAATTTTT 1320

CCATTTTTTA GAAAATGGTT TTCTATAATA GAAATATTAT ATTTTTTTCT CTATGTCTTG 1380

TAATACCATT AATAATATTG ATCAAAATAT TAAAACTTTC TATTGACCAT ATTTCTGATT 1440

GAACTGGAAT TATAATATAG TTTGCAGTAT TTAAAGCGCT CTTTAAAAGA AAATTTTGAA 1500

TCGGAGAAAA TGTTTACCAA AATATAATCA AAATAGCAAT TGTATTTGGA AAGAGATATA 1560

AATATTCTAC ATATTTATTT TTCATAAAAA CTTTTTTTTA AGATAAGATT AATACCAATG 1620

CTATGAGATT TATTTTTATC ATTTTAATTT TCTTTTAATA AAAAAACTTA ATAAGTTATC 1680

AACTTAGGTC AACCACAGTA AAACAATTAA AATGACATTT AAAAAGATTG AAATTTAGTA 1740

AACAAAAAAA AGCCCTTATG GGCATCCTTT TCCGGTTTAG GATAAAGAGA AATCTTTATC 1800

TTATACATTA AGAATAATAT ATTATAAAAA AATAATCAAT GGATTATTTT TTTTATTCTT 1860

CTTTATTTTC ATACTCCATC ACAAGACTAA ACAAGATATC CCTTCTATCC TTAATAATTT 1920

TTTCCAAAAT AAAAACTGCT CTTTTGGTAT CTCTTATGCA AAAAGAATAA ACTTCCTTAT 1980

CTTTTATTAA AAATTTAACA GGGGGCATAC ATGAATTATC ACAATCTTTT TTTCTATCTT 2040

TTTTATGTTC AGATGATTCT AATTGATTTT TCAATATTTT TCTGTATACA GATGAAAAAC 2100

CCkGTTTTTT AATCTCCTCA ACAGAAATAT TTCCCTCTAA GACTTCTTGA TAAATTTTTA 2160

TATATATATA CGCTTGGCTT TTATTAATTA CATAAGACTT AATAAACTGT TCGAAACTGT 2220

CGAACCCGTC ATATTTATAA AGATTTTTTT GTTTGATTTC GTATAAAATT TTCATTCTTT 2280

GAAATTTACT ATCAATATCT AATTTTAAAT TGTATGCTAG CTATTCTTTT AATTCATTGT 2340

AATTTTTTAA TTCCCTGTTC TGATTGATAT CCATTTCTTC TATTTGAGTT TCTACCCTTT 2400

TATACAAAAT TATGTCTTTT TTGTTTTCTT TTTTTTCCAT TTTTACTCCC TTTCTTTATT 2460

TTTAGAAAAA GACCATTCGT ACACTAACAC GATTTTCTCT ATAAGATAGA TTACGGTTAA 2520

AACAATTTTC ATTATAAAGA AAAAATATTT TTCAAAGTTT CTTTAATTTC TTGATAATAT 2580

ATTTCTCTTG AAGAAGGTTC GAGTCTTTTA GTTATAAGAA CTTTTATACT ATTATAAAAA 2640

TGTATCTTGC CCTTGATATA TTCTTTGTAT TCTTTATGAA TCAGATTTTC TACTTCTTTA 2700

AGAATATTTC TATTTTTTAT AAATTGGTTT TCTACAATAG AAATATTATA TTTTTTTTCT 2760

CTATGTTTTG TAATGCCATT AATAATATTG ATCAAAATAT TAAAACTTTC TATTGACCAT 2820

ATTTCTGATT GAACTGGAAT TATAATATAA TTTGAAGTAT TTAAAGCACT CTTTAAAAGC 2880

AAATTTGAAC CTGGAGAAGT GTCTAACAAA ATATAATCAA AATCGCAATT TATTATATTT 2940

CTATTTAAAT AATATTCCAA GAAAGTTTCT TGATCGGTTT CTGTATTAAA TTTTTCTAAC 3000

ATAGGATGAG AAGGAATTAT ATACATATAA TCATTAATTT TATTTAAGTA TTTTTTAAAA 3060

TAGAAATCTC CTTTTAACAT ACTATACACA TTATATGTTT CGGCGTCAGG AATATACTTG 3120

GTAAAATATG ATGTTAAAGA ATTCTGTGGA TCCAAATCAA TTAGTAATAT TTTTTTACCC 3180

AATTCCTTCA ACAAATAAGA AAAAAGTATA GAAAGTGTGC TTTTTCCAAC TCCTCCTTTA 3240

ACTGATGTGA GTGCTATAAT ATCTGGTTTT TTAATATCCA TTTATCTAAA ATTCCTCCAT 3300

TTGTTAGTTT TTTGTTGTAA AATTTATATA CTTTTCTTTC CATTTTTTTT AAATGTTTTA 3360

AACTATACTT GTAGTATTTG GTGTCTTCTT TTTGTCTTTT TCTTAGCAAG GTTCTTAGAG 3420

ATAATAAGTA ACACTTAATA GATCCGGTTT TAAATATAAA TTCTATATAA TATAGTTTTT 3480

TTATTGCGTA ATTTCTATTT TTTCCTATTT TATGAAGAAA TGGTTTTTCA AGTCGATCGT 3540

ATCCATACCT TATTCCTAAA AATTTATTTT TTCCTTCTAT TGGGAAAAGA TTGAAAGAAT 3600

ACTTATCTTT ACTATTAAAT TTTTTAAATT CTAGTTTCAG TCTTTGTTTT TTGCAGTTAT 3660

TTCTAAAATT AACTAGATGG TAGAATATTT TTGTGTAATA TATTTTCTTT CCATTCTTTT 3720

CTTCTATTTT GTTAAAAATA TTTTTTATTG GATTTTCATC AATATTCATT AACGCTTTCT 3780

TTTGTTTTAG TTTTCTAAAA ATTGAATTCA AATCTTTTTC CTTATTACAT TAATCAAATA 3840

ATTGTTTAAT GTTTTATTAC TTGTAATGTA AATATGTAGC TTGTTTAAAG TAAAATAATT 3900

AAAGTTCTAG TTGTAAAAAA GTATTGTGGA TAAGAAAATG GATTTCGTCA ATTTACAAAA 3960

GGTATATTAA CTGATTTAGA TAAAAGTCAA AAATATTGTT TTGATTTATA TCAGAATTGG 4020

TAGATTGCkA TGTTTTAAAG TAAGTATTTA GAATAGCTTT TACTATTAAG CTTGCrkACA 4080

AAATGGTGTT GTCAGCTTTA TTCCAATCTT AATTTCATTT TATCAAATCA AATTAAGATT 4140

GGAATCAAAT GAAACAAAAA AAGTGAATAA GAATTCATCA AAAGAAATTC GCAAATATGT 4200

TAAAGGATTA CTTAATATTG TTAAAAATAT TGTGCCTGTT ACAACTAAAA GGTTGAAGAA 4260

CTATATTTAA AATTGTGCAG GGTATCTAAT GAA 4293

(2) INFORMATION FOR SEQ ID NO: 30:

(i) SEQUENCE CHARACTERISTICS:
(A) LENGTH: 4228 base pairs
(B) TYPE: nucleic acid
(C) STRANDEDNESS: double
(D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 30:

TTGGAAAGGG AAAnGCAAAT TCTCAACAAn ATTATCAAAA TTTAAnGGAA AATATTTTCA 60

ATATCCTTAT TGAACAACTA AAAAAAGAnA CAAATATTGA AATTCTAAAG CCATTATCAA 120

AAAATATTTG AATAACCAAA AGAAAATAGA ATACAATAAA GTATTTGGTA TATATCATCT 180

TGAATTATCA GAAATAATAA AAAATGAAAA AAATTCTTTA ACCACAGAAG AATTTAGCAT 240

AAAGGnCCGT ATGAGGATTT AACATGGAAA ATGCmCCmGA mCCTATTGAA aCTGTAAAAA 300

aGGGTAAATG TAAGGTTGAA TGCCAAAATA AAGAACGyTT TATTTTGATT GAAAAAGAAA 360

ATGGTAAAGC AATGTACCAT ACAAAAATAA TGATGGAyAT TTATAAATTT GGAGTTTATG 420

AGAAAAAACA CGAATTTAGA TTATCATTGA GGGCCTTATT TAAyGGGGAA AGAATTGTTG 480

AAGAAACTCA TTTrTACCCA ATTAAAGAAG GAGATAAGTT yATTGGTATT TTTTATGGCT 540

ACAGAAAACC AATTAAAAAG CCGTTAATAA AGTATCAAAT AAAyGGGACT ArAAAAGCAT 600

ATGCATTAGC AAGGGCATAT TATATGGAAT TTAGATTTAA AGCAGGAAGT GTTTTTTGCT 660

ATTTyAAgGG mTaTATCGAT TATTAGATAA AAAAAGAACA AATAATCACT ACAACAAAGT 720

TTTATTTAGT ATGTTTACAG ATTTAGAACA ACAAGTATAT AAATTTTATG GGAAaAAATA 780

CCCGGAGCAA GGACCGTTAA TAAAATGGAT ACTAAAAAAC CTAAAATAAT AACAATAGCG 840

TCAATTAAGG GCGGTGTTGG CAAAAGCACA AGTGCAATAA TTTTAGCAAC GCTATTATCA 900

AAAGACAACA AAGTACTTTT AATTGATATG GATACACAGG CTTCAGTTAC TAGTTATTTT 960

TATAAAACAT TAGTAGAAAG TGAATTTGAT TTACTTGAAA AAAATATATA TGAAGTTTTA 1020

AAAGGAAATC AATTAATAAA TGATGCAATT ATCAATGTTG ATCATAATTT TGATTTGTTG 1080

CCAAGTTACT TAAGCTTGCA CACTTTTAGT GAAGAGCCCT TGCCTTATAA GGAACATAGG 1140

TTAAAGGATA GCTTTAAATA TTTAAAATTT AAATATAATT TTATTATACT TGATACTAAT 1200

CCCCATTTAG ATTCTACGTT ATCCAATGCT TTAGTTGTTA GTAAACATGT TATAGTTCCA 1260

ATGACTGCAG AAAAGTGGAC TATTGAGAGT TTGCAACTAT TAGAGTTTTT TACGGATAAA 1320

TTAAAGTTAA AACCCAAAGT ATTTTTATTT GTAACAAAAT TTAAAAAAAA TAAAACTCAT 1380

AAAGATTTAT TAGAAATGTT GCAAAAAAAA GAAAAGTTTT TGGGGATAAT ATCAGAACGT 1440

GAGGATTTAA ATAGGAGAAT AGCAAAAAAT GATAGATTTG ATTTAGATAA AGATTATATA 1500

AAGGAGTATG TAAACGTTTT AAATAATTTT ATTTTGAAAA TATGAAATTT GTCCGATAGT 1560

TGGATGAATT TTTTTAACAA AAAGGATAGG AGATTAATTA TGGACGTGGG AATAAAAATA 1620

AACGATAGGG TAATATCAAA AAAGGAAATA AAAAAAGAAT TAAGCAATAA AGATGAAATA 1680

TTAAAGCATT ATAATTTGTT GAAGGAGCGC TTGAAATCTA ATTTTCAAAA AGAAATCTAT 1740

AATAAGATAG AGAGTATGAA AATTTTAAAA GAAATAAAAG ATAATGAATA CTATAAACTT 1800

GATGGTTATA AAAGTTTTGA TGCTTTTATA AAAGATTATA AGTTAGCCAA AAGTCAAACT 1860

TATGAATATT TGAAGATAGC ATCAGCTATA GAAAATGGCG TAATAGAAGA ACTTTTTTTA 1920

TTAGAAAATG GAATTAAAGA AACTATAATC TTTTTAAGAA ATAGTAATTC AGATACGGTT 1980

AAAAAATCAA AACAAAATCC AATAAAACCA TTAAGATTTC AACTTAAAAG CaAAGAAAGT 2040

TATGATTTTT ACAAAAGTAA TGCTAAATTC ACGGGATTTT TATTAGATGA ACTTTTTGAA 2100

AGTCAAAAAG ATTTGATTAA TAAATTCTTA AGAAGATATA AGCAATTAAA AGGATAGTAA 2160

AGGTATTTTA TGACTAATTT AGCGTACAAA ACGTATAACA TAGAAAGCAT AAAAAATGAG 2220

TTTTTAAACA TAGGATTTAG TGAAGAGGCA ATAGATTTTG TTTTGCTTCA TAATGAAAAT 2280

TACAGCTTTG AGGTTTTAAA AGAAAAATTG ATTAATGTAG AGAAGAATTT GCAAAAAGAT 2340

ATATCTAGTT TGGACATTAA GATAGATACT GTAGAAAAGA ATTTACAAAA GGATATATCT 2400

AATTTAGACA TTAAGATTGA TGCCGTGGAA AAGAATTTAC AAAAGGATAT ATCTAATTTA 2460

GACATCAAGA TTGATAATGT AGAAAAGAAT TTGAAAAAAG ATATATCTAG TTTAGATACT 2520

AAGATTGATG TTGTAGAGAA GAATTTAAAT CTAAAAATAG ATTTTGTAGA AAAGAGTTTA 2580

AATGCCAAAA TAGATAGTTT AGACGTTAAA ATAGATAATG TAAATAATAA AGTAGATTAT 2640

ATTAAAAGTG AACTTATTGc CAAGATAGAT AGTGTAGAAA AAGGGTTAAA CGAAAAACTT 2700

AATACaGGAA ATAGGCTAAT ACATTTTATG ATATTAACAG CAGCGATTCT AGGCCCAGTT 2760

TTAAATGCCC TATTTATGAG ATATTTACAA TATATCAAAT AATGATGTAA TGCATAATTT 2820

GCTTTTTTCA AATAGTTTAT TATCAATTAA AGCTTATTTA AGCTTTTAAA TAAAGTAACT 2880

TAAATAAGTT CTTTTATTTT AATAAATACA AATTGATTTT AATTCTAAAT TGAACTGAAT 2940

TTAATTGTTT AGTGAGTTCA CCTAAAATAA ATTAAGCTAA GCCCGCGGCT TTATTAAGCT 3000

CTTTAACATG AGAATTTAAT AAAGCTTTTA TTTATTATAA TAATTTCTGT AAAAAGCCTG 3060

ACAAAAATAG TTTTTGTTAT ACATATGTAT ATGTATAGCT AAAAAAATAT ATTGCTATCA 3120

AAAAAATCCA ATTAAGTTGG GTTTAGCTAA GTTCTTTAAC AAGAGAATTT ArmTAAGCCC 3180

tATTTTTTTG TAAAATTTTT TGTAAAAAAG TTGGCAAAAA TAGTTTTTGC TATATACTTA 3240

TTTtTAtAAA TAAcaaGGAG kAAAAAGATG GAaAATCTTT CAAACAATAA TAATCCACAA 3300

GAAAATATTC AAGGAGAGCT CAAAATGATA AGTGTTAATC AACAAAGTTT TACTGGTTGT 3360

GAAATAATTG AGGAAAAATC TTCTCCCATT AAAGAAAAAA GTAAATTAAG TAAGATAGGT 3420

AAGAAATTGC CAGGAATAAG CAGTCAAGAA TGTTTTAGAT TTAATCGAAA TATTGATTTT 3480

AGTGTGCAAA GAAACAAsTT aGATAAATAC GGTGCTAGTG AAGTAGGCAA TATTCTTGTT 3540

GGAGGTGCTG GrCTkAAAGA TTTAATGATA AACAGAGTGC TTAAATATTT TrrwATGAGy 3600

CTACCTTTTG AAGAGAATTT rTATATGCTC AAGGGCAAAG ArTTAGAGAA TTTAGGATTT 3660

AGAGArTTTG TTAAAGCACA yrGTGATAAT ATTrATrTTT TGTATAAAAA CAAATATGCC 3720

AAyGGwGTTG ATAAGTATAA yTATTTCAAA AAAATGGGyA GTTCAsAAAC TTTAGTGGGC 3780

TCAACAATTG ATGGCTGGTT TATTAATAAT AATGGCGATT TAGAACTATT AGAGATTAAA 3840

AGTAGCGACT CTCATTATAT GAGTAGTGCT ATTGCTGAGT ACAATAAAAA TGGCAATTTT 3900

TTAAGCAGTA AATATTTTTT CAAATATTAT GTACAAGCAC AAATGCAGCT AGCATGCACT 3960

GGGCTTGAGT ATTGTAATTT GTTCTTTTTA ATAGATGCTG CACCAATTAA CTGTAAGATT 4020

AAAAGAGATG AGGCCTTAAT ATCAAAAGTG TTTGAATTTG TTAATAAATG TGAATTAGAA 4080

ATTATAAATT TAAAAAAAGA TATTTATAGT AACTATAGAG ACGATTACTT AATGGCACAT 4140

AATTTTAATG AGGATACGTT TATAAAACTT GTTGAAGATT TAGTAGAAAG GAGTGATTTT 4200

TATAGTTCTG GAGTTGAGTT TGATTGGG 4228

(2) INFORMATION FOR SEQ ID NO: 31:

(i) SEQUENCE CHARACTERISTICS:
(A) LENGTH: 4137 base pairs
(B) TYPE: nucleic acid
(C) STRANDEDNESS: double
(D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 31:

TTATTTTGAA TTAAAACTAA TGTTTAGTAA TTCAAATATA TAAGGAGAGC ACATTTTGAA 60

AAACCTAAAT TAAATATTAT TAAGCTTAAC TTTATTACAG CAATACTGAA TTCAATTTTC 120

ATATCATTTT TACCTATTGG AAAGGTCGAT CCAAAACCCG ATATCAATAC TAATCCAGAA 180

AATATCCAAA ATTAAAGTTT AGAGAAGCTT TTCGCATAAA TATTTTATAA TTTTTAGGAT 240

TATATCGGCA ACTATGTTAA ATATTACTCG AATGGCTGTA CTTTGATATT AAAATCTGTT 300

ATTTGTGGAG TTGGTATTTC TGATTTTTAA ATAATTTCAG GAGTAACTAT TTCTTTGAAA 360

ATTTCAGGAA TTGTTGTTTT AGGAATGATC ACTTTTTCAG TAATTTCAGA AATAATTTCA 420

CCAAGGCAAT TTTATTCTTT GCAAGAATTT TGTTCACAAT TGTACAACAT AATCAAAAAA 480

TTAAAAAAGT AAAATGCAGT ATGAAAACTG ATAGTGAGAT TTCTTATTCA AATAGGAACA 540

TCTACAATTC TATCCCAAAA GTAAAATAGG AAAGAGTTTG TTAGAAAGAA AACCGTATTA 600

ATTTCTTGGT TAAGTTATGT AATAGTAATG TATTGTTATT AATTTATTAT ATAAAATTTA 660

GGCATAAAAA ATAGGAGGTA TCATCCAAGC ACACTTTAGG TGCTAATGAG AAAATAAAAA 720

TTTTTTAAGT GGAGAATAAA GAGTAATCTT TGGGCAAATC GAACAAAGAT TGCTTTACTG 780

TCTTATTTAA CTATAGTAAG TTTATCTCTA AAATAAGATT ATCCAAGCTT ATTTTAAGCT 840

TTTATTCATA TTTCTCAGCA AAATCAATTT AGAAATCTTT TAATTTTTAA TTCTTGTATA 900

ATGCTTACAA AAAATCATAA GTATAGAATC CGCACAAATA CCAAATGTTT TTTTATTCAT 960

CATAACTTGA TCCGTAAATT TACGAACACC AATAATACGG TATCGTAAAA TAAATAACTT 1020

TATCAGCCTC AATAATAACT AAGAATACAA ACTTGGCATT GCTAGTAAAT TTTAGCAAGA 1080

TGCTAAAGGA TGATGTGCCT GGTTATAAGG ACAACCAACA ATACTTTAGA GAATGCAATT 1140

GTAAAGTATA AGGAAGCCAT AAGTAAGACT ATTTGGGGCT AAGAGTCAAA TTAAAATAAA 1200

GACAGAAAAC AATAAGTAAG AGAAAAAGAA TGAAAGCTTA ACTCATTTAA GAATAGTTAG 1260

AGGTGTTCTT TCTGTTGTTA AAAAACCACT GAAACAGCTT TGCCTAGCTT GCGCTGATTT 1320

TATTGCTACT GCTGCCAGTC TATCTTGTAG TGAATTTAGG CAAACTGCTG AAGATTTTAG 1380

TGTTTGCTGC TAAGGAGTAT GCTAATGGAA AAGGAAAAAT AATGATTTTG ATGTTATTAG 1440

GTGCTATTTC TAGTATGGCC TATAATGAAT TTGAATAGGA GGTTTCAAAG AGTAAAGACT 1500

TTTGCTAATA ATGAAGGGGG CAAATTAAAC AAGATGACTG CTACTATTGA TAAGTTATGT 1560

GCTGTTTATA AAAAAGTTAT ATCTTAAATA CCTTAAGTTT TTGGCCAACT CTTTTCTCTC 1620

TTTAAAACAC TTTGCCTCTA TGCTTGCTTT ATAATAAAAT AATACTTGGA TAATGAATAA 1680

CTAAAAAAAT AAGGAGGTAT TAATGAAAAG GAAAAGCAAT ATATGTATTT CACTTCTAGT 1740

CACAATATTA TTTGTGTCTT GCAAGTTTTT TGGAAATAAA AGCGCAAGTA AAGAAAAAGA 1800

AGAAACTTCT TTTTCTGATA CTGCTAGCAA GATTAGTAAG TCGGGAACAG CTGCTTCTTC 1860

AGACAAACAA GAAAAAAATA CAAGTGATGT TACAGGTGAC GCCAAAAAGC ATACTAGTAG 1920

CCCTTACATG CTTGCTGATG CCCTTATTGT TAGTGATACT ACTAATAGAG ATAGAGATAA 1980

GCAAGAAAAT AAAGATAAAT TAAATGAAGA AGATAAAAAA AAGCTTAATG CTTTTTTTAG 2040

CACAACTAAA ACATATCAAT CTAGCCTAGA TTCCATTTAT AACAAATATA CAGGCTATTA 2100

TAATACCATT GATACCTATG GCAGCTGTGA TACGTATCGC ATTGAGTGTT TTAGTGTAGG 2160

ACCTTCTGAA AAACGTAAAC AAGCTCTTGC TGATCTAGAG AAGTTAAAAC TAGACGAAAA 2220

GTACACTCAG CTTAGCACAA TGTTAAAGAG TGCTGTGCCT AGTTATTACA AAAAAAATTT 2280

AGATGATTCT ATTGCACAGT ATAAGGAAGC CATAAAGCAG GCTATTGAAG CTGAAAGTAA 2340

AATAGAGACA GTAAAAGACT ATGCAACAGC TCAAAGTGCT GCCGATGACG AAAAGAAAAG 2400

AAATATAGAT AATTTAAAAA TAGTTAGAGA TGTTCTTCTT ATTATTAAAA AAACTATTGA 2460

GAAAGCCAGC CGATCTTATG CTGATGCTTT TGCTATTGCA ACATCTAGCT TATCTTGTAG 2520

CGAATTTAAG CAAGCTGTTA AAGAGTTTAA TGATGCTGCT AAACAATATG CTAATGGAAA 2580

TAAAGGAGAC AATGCTGTCA ATGTTATTGT AGGCACTATT TCTAGTATGC CTTATGTCAA 2640

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GAATAAAATA AAATTAGGGT TTTACACCAA AAACTATGTT CTGTAATATT CAATATAATT 2820

TTCTTTATCT ATGATAGAAG AGTCTTTAAA CAAAACATTA ATCTTATAAA ATTCTTCTCT 2880

CCTCTCACAT ACTTTCACAT CCTATTTAAA ATTAATACTA ATGCCTCCCA AGAAGCCCAT 2940

TATTGCCATA TGAATTTAAC TAAAATACTA TCGAAAAAGA AGATATGAGC TACAAAACTC 3000

TTTAAAAAGC CCCATATCCA AACTAATCAA TTTAATCACT TTATGTCTGT CTTACTTAAT 3060

AAATCACTAG AATCAACTTA TCAAAAGTAC TTTCTTACCC CCCCCAATCA TCGTAGTTAT 3120

CTTTTTAAAT CATATATGAT ATATGTATTA TTATCAGCTT ATCTTCCCTA TATATTAGGT 3180

AAAAAATAGA AACAAACAAC CGTGCATTTG CGACTCAATA AAACACCTAT CAAAAACCAA 3240

TATTCTACTT TCATTATACA GAATAAAGCA ATATAATATA TTTAATATAT ATTATCGCTT 3300

AAATTTATTT AAACTTAATA AATAAATTAA GGGGAGAATA AAAAAAGAAT GATAAAAGGC 3360

AATACGTTTA TTTTAATATT AGTAACAACA ATGTTTGTAT CATGCAAATT CTATGGAAGT 3420

GATGATACAA ATAAAAAAAA TACATCTCTA AATGGTGATA CTAGAGAGAT TGACAATATA 3480

GGTTCAGTCA TTTTAGAACA AGACGGAAAC AAAAAAGGTG ATACTACTGC TAGTAAAGTT 3540

GCTTTGGATC AAGTTACAGA ACATGCTAAT AGTGAACTTA TGCTTAATGA TGACCCTGAT 3600

TCTAGTATTA GTAAATACAA CCAAGAAAAT ACTACTGGCA AATTAACCGA AGAAGATATG 3660

GATAAGCTTA AGGCTTTTTT TGTAAAAACT ATAACATATC AAGGGATACT CAATTCTATT 3720

TATAACAAAT ATACACGCTC TTATAATACC ATTGCAACTT ATTCTGGTTG TGCCAATTAT 3780

AATAGTATTG GATGTTTTAG CGAGGGCCCT TCTGCAAGGC GTAGTCAAGC CCTTAACGAC 3840

CTAGAAAAAA ATAAACTAGA TGAAGAGTAT ACTAAACTTA ACCAAATGCT AAAGGAGACT 3900

ACACAAGATT ATTGCCCCAA AGCTCTAGAC AATGCAATTG AGGAATACAA GAGGGCTATA 3960

ACAATAGCTA AAGAAGCTGA AGATAAAATA AAAAAAATAA CAAGCTTTAC AATAGATGAA 4020

GGCAATAATA ATGAGGAAAG AAAAGAAAAT GTAGATAACT TAAAAAAAGT TAACAATATT 4080

CTTTCTATTT CCGAAAAGAC CATAGAAACA GCTAGTGTGG CTTATGCCAA TGCTTTT 4137

(2) INFORMATION FOR SEQ ID NO: 32:

(i) SEQUENCE CHARACTERISTICS:
(A) LENGTH: 3760 base pairs
(B) TYPE: nucleic acid
(C) STRANDEDNESS: double
(D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 32:

ACTTTTTCAT TTTTGAAGCT TTATATATTt TTATaCAGTG ATTAATTCTA AATTTTTCTT 60

TATTTyaTAA AATTTTTGAT AAATCAATAA TCTACAGCCA ATATTAAAAT CAAAATATAA 120

TAACAAATCT ACAAGTAGAT AAGATGCCTT CCAAATCTTT TTTACTCGTA GACTTGATTA 180

AAAAAAAAGA ATAAATTAAA TGCAATTAAT AATTAAAATT ATAATAAAAA AAATTATAAA 240

ATAATATCAA AGTACAAATT TTATATTTTA GTCGTATAAT TTTGATTCTA AACTATCCTT 300

TATATGATAA AATTTTTGAT AAATCAATAT AGATTCGTCA TAAACAATAT TTATATTTAA 360

ATTATCAACA TTATTGCTAC ACGACAAATA TAAAAGCTTA AAAAAATTTA TATCACTAAT 420

ACTCAAATAG AATATGAAAA CCTCATTTTC AAAAACATTA TTTCCTAAAT TTGTCTTAAT 480

TTCACAAAAT AATTTAATTA ATTCTTTAGC TCTTTTAGAT CCTAAACTAA ACAAGAACTC 540

ATTCAAAATG TCATGACTTT TAATTGGAAA CAATAAAGAT AAAGGGTTTA TATCGGTTTC 600

CATATCTGAT AGTAACTTAT AAAACTCATA AAGACTTTTT TTTATGTCCT TTAAAGTAAT 660

AATTTCAATG GCATTTAGTT CTAAATAAGT GTATATGTTT TCCACCATAT ATTTTTCATA 720

AAAAAACATT TCCATTTTAA ATGGAATACA TTATATTTCT TTTTTAAGAT ATTTTAAACG 780

TATGCTTTTA ATTTTACATT TATTATTAAT GAATTTATTT GAGCTAGCTT TTATATTTGC 840

TAGTAATAAA AAAGGAAGCG AAGTAGATAC TATGATTACA GCTATTGATA AGTTAATTAC 900

TATTTATAAG AAAGTCAAGT CCTAAGGTTT AAAATAAGAG AAATATATAT ATAAAGAATT 960

TTTTGTGCTA TATTTATTTA CAACAAAAAT GAAGATTAAA TTTTTAACTA TAAGTTTAGC 1020

TATCTTGcTA GCTAACTTGA TTATTATTTT ATTAAACTTG GTTTTATTTA TTGTAAATAC 1080

ATCTACTAGT AGCCCTTATA TTGTTCCCTC AGAAAAGATT GATATTTTAC ATCAAAGTAA 1140

TACAGGTGCG GTTAAATTTA AGATTTCCTT AATAAATCAT TTGGGCTCTG TTGCTATTGT 1200

TTATGATTAC AATTCTGCTA GTGAAAGATT TTATTTAGAT TTTGAGATTG TAACAAACAA 1260

GAAACCTTTT AATCTTTTAG ATGTTTCTCT TAATGATGTT GTTATTAAAC CTGAGGTTTT 1320

ACTTGCATCC AATAGCAAAT TAAGATTCGA AGAGGGTCAG TATGTTTTGA ATTTTGACGA 1380

CTCTATAGAA AAAACAGGAT TTTTTGTTGA CTTAGACCTT AGAAATGAAT ATTTAAATCT 1440

AGCAGAAATT GCAAGAATCT CTGGCATTAA TTTTCGTGTT AAATGCATAG AAAGAGAAAC 1500

AGGTGTTTTA CGAAATGTTT TGTTCAAATT AAGTGTCGAA AAGGGTAAAA AATTCTTTGA 1560

CCTTATTGAA AGATATAACA ATAATATTGG CAAGGTTTCT TGATTATTTG ATTTTGTGGG 1620

GAAAAAAAGT TTGATTTAAG GAAATATGAG GAATATTAGC AATtGTATCA AATATATTAT 1680

ATTAACAATG CTTATTGGAT TATTAATTTT TTGTTGTGCA ACCTTTGTTT GGTTGATTGG 1740

AATTTTTTAT TCAAATAACT TTAAAGAAGA GCGGAATTAT TCAATAAGCC CAATAGATAG 1800

TGTTATTATG CGTAAATGTT ATTTTAAAGA ATTTAAGTCT GGACTTATTA AAAGCGTATT 1860

CTTTAAGAAA TTAGATGTAA ATGTTAACTC TAAAAATTTT AAGGAGCTAA ATAAGGTAGA 1920

TAAACAAAAT CTGCTAAATT CTTATCCATC TTATCATATG GAGTTTGTCG TAGTTGATAA 1980

TGGATTTTTA ATGAATTTTA AAAATGTTAT TTTTAATGGT ATAGATGATG CTAAATTATA 2040

CGATCAACGT GATATGGTTT ACGGAGGATT TAGATACTCA AAAGAGGCTT ATTTCCAAAT 2100

TATTGGCAAT TATGATGTTA AATTAAATAA AATGAAACAA TATACTCCAG CAATTGTAGT 2160

AAATGTTTTC AAAATTAACA TTAATGATGC TTTATTTAAC TCGTTATTAA AGCAAAAAAC 2220

TTTAAAAGTT ACTTTGATTT CCCATAATAA TAAAGAGTAT ATTTTACAAA CTAATAATTT 2280

CTTATCAAAG TATAATTTTC AAACACCAGA AAAGGAGAAT AGTTCTTACT AAGAAATTAA 2340

AGAATTTTAT TTTAAGTTTA ATAATGGGTT GATCATGTCT TATGGTCAGC AAACCAAATA 2400

TGCCTCCCCT TTATCAGAGC AATTATTCTC TATACCCCCT CTTTTATATA CTTTCTATAT 2460

AATAAATATA GTTTCATAAA ACAATTTTTA TGTAAAAAAA GCTAACTGCC TTATTATTGC 2520

TATATATTTA TAATATTATA TTATATAAAA AGATAGGTCT GAAACAATAT GGATAAATTT 2580

TTAACATCTA ATCATCCTCC AATAATAATC TTTACTATTG GGGCTTTGTG TGCAACCGTG 2640

CTAATATGCT TAATTATAAT ATTTATTATA CATGGCATAA TAAATCCCAT TCTTATAAAG 2700

AAATTCAAAT CCATAAACAA CAGTCTACAA AAAATAACAA AAGAATTTGA AGAGATGAAA 2760

AAAAGAGTAG GACAATTAGA ATCCATATCA AAAAAATTAG AGCTAAATAA TAAACCTGCA 2820

AAAAAACCTA TAAGCTTATA TGGACTAAAA AAATCCCAAA TCTAATGTAG ATTTACAACC 2880

GATTATTTTC ATTTTTTTGA TTTAATAACA AATAATCAAT AAAAATAATT TACAAATAAT 2940

TACATACAAA AAGCTACACG CAACAAGGCA AGGCAAAATA AATTATTAAA AAGCGAGCAA 3000

CAATATCAAG CACCCCTTCC CCTATAACTA TTTTAAAAAA AATTAAAAAG GTTTAACGGT 3060

ATACTTATAA ACCTTTTTAA TAAAAAACAA AATTTATTAT AAACCTCAAT AACAGTAAAT 3120

GTTTAAAATA CATATAACTC ATTTTAGCCC ATCAAAAGAC ATTTTCAACT TTATTTTACA 3180

ATTTTTTAAT AAACTCTACA TTGGGTTGGA CAATGGAATC TAAAAATTCT TTATTCTTTA 3240

ATATTTCATA CTTTTCATAG TATGAAAAGT ATGAAAAAAA TGATTAATAC TACAATACAT 3300

TCCTAAAATT TATTTGTTTT TTTTATGATA TAACAAAAAA ATGTTTTATA AATTTCCAAT 3360

TCGGATATAA AAATTACCAA AGAGAAATTT TTATGAAAAA AGACATATTA TATACTTTGA 3420

ACGTATTTTT ATATATATCA TTACTTTATT CGTGTTCTTG GATGCCATTA AAATCTTCAA 3480

AAATCATTAA AATTAAGAAG TGTTAAAACT AATGAAAAAT CCATTAATTT TAAGGAAGGT 3540

TTACAAAACA TAAAAGATTT CTTCCTGCAA AGTTATTTAC CTACATATAC TTAATAGTAA 3600

TAGCTTAAAA GCGGATATAA ATATTATTGT AAGAAGAAAG TCGACAATTT TGATACTACT 3660

TAAAGGTTTG GaTTATTCTA TTTCTGcTTT GTTTGTTTTA AAGCGATTTA TGGTTTGAGG 3720

CTAATGTTGA TTTTTTTACC TTTTTTTATT TAAATTTATT 3760

(2) INFORMATION FOR SEQ ID NO: 33:

(i) SEQUENCE CHARACTERISTICS:
(A) LENGTH: 3653 base pairs
(B) TYPE: nucleic acid
(C) STRANDEDNESS: double
(D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 33:

CTTGnGTGCC TGCAGGTCGA CTCTAGAGGA TCCCCGTGTT CATAAAATCC TCTCCTTGAA 60

GGTGTTACTT TTAAATTAAG TAAAAGTAAT AAAAATAGAT AAAAATAGTA ATTTATATTG 120

TACCAAAAAC GAAAAATTTT AGTCAAATTT TGTGAGTTCT CATTGCATGA GAAATTTGGG 180

TTGTAGGGAG GCTGTTATAA ATAGAATGGG CATTTTCTGA GGGTGTCGGC TAAGAAAGAC 240

TACATACTTT AGCTAATATA TAGCAAAGAC TTTGAAATTT AATTTGTATG TGTTTTATAG 300

TCTTTTGTAA TGAGTAGTGC ATTTGCAATG GAGAGATTTT GGGGAGTTGT TTAAAATTAC 360

ATTTGCGTTT TGTTAAAATG TAACAGCTGA ATGTAACAAA ATTATATATT TAAATCCTTG 420

AAATATTGCA TTTATTATGT ATTGTGGTAT GATTAGGACT TATGGAGAAA TTTATGAATA 480

AGAAAATGAA AATGTTTATT ATTTGTGCTG TTTTTATACT TATAGGTGCT TGCaAGATTC 540

ATACTTCATA TGATGAGCAA AGCAATGGAG AGGTAAAGGT CAAAAAAATA GAATTCTCTG 600

AATTTACTGT AAAAATTAAA AATAAGAATA ATAGTAATAA CTGGGCAGAC TTAGGAGATT 660

TAGTTGTAAG AAAAGAAAAA GATGGTATTG AAACGGGTTT AAACGCTGGG GGACATTCGG 720

CTACATTCTT TTCATTAGAA GAGGAAGAAA TTAATAACTT TATAAAAGCA ATGACTGAAG 780

GTGGATCATT TAAAACTAGT TTGTATTATG GATATAATGA CGAAGAAAGT GATAAAAATG 840

TCATTAAGAA TAAAGAGATA AAAACAAAGA TAGAAAAAAT TAATGATACT GAATATATTA 900

CATTTTTAGG AGATAAAATT AATAACAGTG CGGGGGGAGA CAAAATAGCT GAATATGCAA 960

TATCACTAGA AGAGCTTAAA AGAAATTTAA AATAGAAGTT AGAAATATAG GAGAGAGCGT 1020

ATATGAATAA AAAAACAATT ATTATTTGTG CAGTTTTTGC GCTGATACTT TCTTGTAAGA 1080

ATTATGCAAT TAAAGATTTA GAACAAAATG CAAAAGGGAA AATTAAAGGA TTTATAGATA 1140

AGGCTTTGGA TCCAGCAAAA GATAAAATTA CTTCAAGTAG TTCAAAAGTA GATGAATTAG 1200

CAAGAAAATT ACAAGAAGAA GATAAAATAA AGGGTGTAGA AGAAAACAAT AAAGATGAAT 1260

TAATGCAGGG TGATGATCCT AATAGTGGTG TAATAAATTC GTCACCAGTA TTGCCAGAAA 1320

ATAGTCAAGA TAATACACCA ATATTAAAAG CAGCGGAACA AAGTGATGGT CAACAAGAAG 1380

AGAAAGTGAA AAAAGTAGAA GAATCCGAAG CTAAAGTTGA GGGAAAAGAA GAAAAACAAG 1440

AGAATACAGA AGAACGAAAC AAACAAGAAT TAGCTAAACA AGAAGAAGAA CAACAAAAAC 1500

GAAAAGCAGA ACAAGAAAAA CAAAAAAGAG AAGAAGAGCA AGAAAGACAA AAAAGAGAAG 1560

AAGAGCAAGA AAGAAAAGCT AAGGCAGAAA AAGAAGCTAA AGAAAAAGCA GAAAGACAAA 1620

AACAAGAAGA ACAACAAAAA CGAAAAGCAG AAAAAGAAAG AGAAGAACAA CGAAAAGACG 1680

CAGAAAAAAG GCAaGTTGAT AACGAAATTA GAACACTTAC AGGCAAAATA GATGAAATCA 1740

ATAGAAATAT TGATGTTATA AAAGAGCAAA CTAGTGTGGG GGCACAAGGT GTTATAGATA 1800

GAATTACAGG GCCTGTATAT GATGATTTTA CTGATGGGAA TAAAGCTATA TACAAAACTT 1860

GGGGGGATTT GGAAGATGAT AACGACGAAG GATTAGGAAA GCTATTAAAA GAATTGAGTG 1920

ATACTAGACA TAATTTAAGA ACCAAATTAA ATGAGGGTAA TAAAGCATAT ATTATTGATA 1980

CTAGAAGCAC TGAACCCCAA TTAAAAGAAA ATGTAAGTGT TAGCGAAATT AAATCAGACT 2040

TAGATGAACT AAAATCAAAA TTAGAAGAAG TTAAAGAATA TCTTGAAGAT AAAGATAATT 2100

TTGAAGAAAT TAAAGAATAC GTTGCTGGTA GTGAGGATAA TTATGATGAA GAAGATTAAT 2160

TTTAGATATA ACTAAATTTT GTATACACAA AATAACAACT AGTAAAAAAA TGACTAGTTG 2220

TTATTTTTTT GTAGATTTCA TTGTTATAAA TATAGAAATG TTTTCTATCA AAACTTTCAT 2280

TCAAAAAATG CCAAAAACTA TTGCTCAAAA TATTGTTTAT TTATATACTC TCTAGAGTTA 2340

TGATGAATAT AAATGAGATT TCAGATTTTT ATGATAATTT ATATAAGAAA ACAAAAAAAG 2400

AAATAGATAA ACTTATAAAC AAGCTCTATT TAACTAGCCA AATAACTCTA AAGCAAAAAA 2460

GACAAATATA CAGTGCTGTT GAAAAAATGC AAAAGTACGT AATAAAAACC GGAAAAAGTG 2520

TTTTTTTAGA ATCGGAAAAA GAATTTGTTA AAGACACTTT GAAAAGAAAA AATCTAACAA 2580

AAAAATTTCA AAGTTTCAAA GTTGATTTTA GCTACAAGGA AGGAATGCTA GAAAAATGTT 2640

TAGAAAGATT AGGAGAAGAT AAATCTATCG AATTTTTGAT TTTTGTTTGC CAAATTCTTA 2700

ATGGGATAAG AGAAAAAGTA TCAGAATTAG ACTTTCAAAT AGATGCGATT AAAGAATTTA 2760

GAGATATTTT ATTTTTGAGT ATACACTATT ATGATAAAAG ACTTTTCACC AGTAAGAATC 2820

TTATGAATGA AATGAAATAC TTTTTCGAAA AAGTAGAGTT AATTTATAGT TATATGCAAT 2880

AAATTAGTGA ACTGCTATTT CTAAATCAAA AATTATAGAA ATAGCAGCGA ACTAAATCAA 2940

TAAAAGCTAA CAGATATTCC CTGTTAAATA TCAAGAAGTT ATCAGTTTAT GTTAACAATT 3000

AACAAATTGC TTTACTATTT AGAGTAACAA ATTGTTACTT TTGTTATTTT AGAGGATTTT 3060

TTTGAAAAAA GTTAAAAGAT CTTTTGATGA TTATGTTGCA TATTTTAGAG AAGGATCGTT 3120

AAGTGATGTA GAAATAGCGA AGAAATTAGG AGTTTCTAAA GTAAATGTGT GGAGAATGAG 3180

ACAAAAGTGG GAAAGTGGAG AAAGTGTTGT TAACGGGGAC TCTAGAGTAA CAATTAGTGA 3240

AGATACTTTT GAACACCTTT TGTCGCAAAC CTTTAGATCA GAAGTTAACG CTAGGAAAGT 3300

TAGAAGCGAA TTGGATGTAG AGCGGTCTAA TTTAGAATTA GGATTTATAA ATGCATTTAA 3360

GCAATATTCT AGTGTTGAGC TTTTTAGTAT GCATACTAAA ATAGAAAATT TAAGAGCCGA 3420

AATTGACGCT TTAAATAAAG CAAGTAGTAA AAAAAACAAG CAAGTTGTTA ATGGAGAAAT 3480

TAATTCTTTA AAAAGCGAGC TTGATGAATA TATAAAAGAG TGTTCAATAA GAGAAATGGA 3540

GCTTTACTAT GAATGTATGA AAAAACTTGC TACGGTTAAT GGAGCTGAAA GCAAAAGTAA 3600

CTACAAAAAT AGTAAAGGGC ACAAGTGAAC TTATATCAAA CAAAACTTTT TAC 3653

(2) INFORMATION FOR SEQ ID NO: 34:

(i) SEQUENCE CHARACTERISTICS:
(A) LENGTH: 3432 base pairs
(B) TYPE: nucleic acid
(C) STRANDEDNESS: double
(D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 34:

CAnTTTTATG GAGGTTCTTA GATTTTTTnG TATTGCTTCT CATAATTAAA AAAATAGAGA 60

ATAAAAAGGT CTTCCCGACT TTTCTTAATA TGGAACAATT GnCAATAAGG AATTCATAAG 120

GAAAACTTAA AGCTAATATA TCATCTATAA ATAATAAAAA ATATTAAAAA ATAGAGGTTG 180

AAATTTTTAC ATATTTTATA CATAATAATA ATTATTACAC GAACAAATAG GAGAAAAGAT 240

GGCATTAAAT TTGCTTAACC ATAACCACAA CAAAACCAAT ACCAACAAAC TACTTTCAAA 300

AGATTCAAGG CTCAAAAAAA TTATTTCAGT AATTAATTAC TTAAATAAAA GTTTTGAAAA 360

AAAATATGAT ATTTCAATAT ATAGAATTCA TTTTAATTCT GAAAAACTAA AAGAGCTTTA 420

TCCTCATCAT CAAATAGATA TCCTTAGAGT TTTAAACTCT AATATAAGCA AAGAAGGCTA 480

TAAACCAACT GTAATAAGAA CCTTAAGAGA AGACCTAAGA TTTTTAATTC ACATAAAAGC 540

AATTGAAAAA AAAATATTAA CATTCTCAAA TAACTTAGGA AAATTTAAGG GAAAATTATG 600

TATATATAAG GTGTCGCCTA TTGCATATAA ATTAATATCT GCTTATTTTG AAGCTTATAA 660

AGCAGACCTC TATAGAAAAA TAAAAAAGAG TAAAGACGGG CTTGATACAC AAAATGTAAC 720

TAAAAATGTT ACTGTATATA TAAATTATCA TAAAAATATA TATAATAAGA ATTCTATTGA 780

AACCGTCTTT AAAAAATCCT ACACTAAGAA AAAAACAAAA AAGAAAAACA AAAAAGAATT 840

CACAAAAAAT AATTTGGAAA AAAGGTTGAA ATTACCTGAA GAAATAACTA AAGAAATTAT 900

AAGTATAGCA AAAAAAACTA AAAATCCAGA TAAAACTTAC AAAAATACGC TTTTTAATTA 960

CAAAGATTTC TTAAACTATT TATCATATGA CTATAAAAAA GAAGATATTT CATATTTCTT 1020

TTTGAGCAAA CTTAAAGAAT ATAAAAATAA AATTCACTTT ATGAGAAAAT ACGCTCCTTA 1080

TAAAACAGAC TTTTACCTAC TTGCAGGAGA ATTCAAAGAT TCGTACCATT CTAAATGGAA 1140

AACAAATAAA AAAACCAATT TTAGCGGACA TGTAAAAGAA ATAGCCAACA ACATTCTAAG 1200

TAAAATTTTA GAAAAGGAGC TAAAGTTTGA ATGATTTACT AGAAAAACTA AAAGCGAGAA 1260

AAAAAGAAAT AATAAGCAAG AAAGAAGCGG AATATAACAA TAATATAAAT AAAGGAACAA 1320

AAGAAAGAAC TGCCTTTTTT AGAATTGAAG AAATAGATAA TAAAAAAATA TATTATACAA 1380

AAATCTTTAA ATATTTAGTA AAATTTAGAA TTGCCAATAA AGACAATAAG CTAAGTTTAA 1440

CTTTCCAAAA ACTTAATAAT AAAAAAAATT ATTATTTATT TAATCTTTTT CCTCTAAAAG 1500

AAGACAACAA GTTTCTAGGA ATAAAATATG GATGGGACAA ATTAGAAAAG CCTTTTTTTT 1560

TAAGGCAAAA TAATAAATCT TATGTAATAA AAAAACTTTA TTATTTAGAA TTTAAATTTA 1620

GCAAAGGGTC TATTAAGTGT TACGTTCAGT CTCTTAGAAC ACTATTAAGA AAAAAAGACA 1680

AAGAAAGTAC CAAATATTAT AAGTTTAATT TAGAACACAT AAAAAGAATG GAAAATACCG 1740

TATATAAATT TTACAGTAAA AAGCTAAAAA ATAAAGGAGT AATATATAAA TGGATAGAAA 1800

AAAATCAAAT ATTATAACAA TAGCAAATCT TAAGGGAGGT GTAGGCAAGA GCACACTATC 1860

AATACTGTTT TCTTATGTAT TAAAGGATTT GGGTAAAAAA GTATTGCTTA TTGATATGGA 1920

TTCACAAAAT GCTTTAACTT CATATTTTAG AAAATATGTT TTTAATTTTG ATAAAAATAA 1980

CATTTATAAT TTATTAATAG GCAATGTTTA TTTTGATCAA TGTATAAGTA AAATCAACGA 2040

TAATATTTTT ATAATTCCAT CACATCCTTT TCTTGATGAA TTTAATGATA AAAATTTGGA 2100

TAATAAGGAA AATTTATTGA GTTTTTGTTT AGACAAAAAT GTTTTAGGCC ATGATTTTGA 2160

TTATATTTTT CTTGATACTC CCCCTAGTTT TAGTTTTATT TTAAAAAATG CATTAAATAC 2220

TACAAATCAC ATTATTATTC CAGTTCAACC TGAAACATGG TCAATAGAAA GTTTGGAGAT 2280

ATTAATGAAA AAAATTACAG ATAAAAGCTA CAATATTTCT ATTGTTGTAA ATCAATTTAT 2340

TAAAAACAGG AATATACTTA AAGAGGTTGA AGATGCTCTA TATAAACGAT ATAGTAACTA 2400

TATAAAAGGT AAAATTCATT ATTATAATAG TATAAAGGTT TTTATAATTA ACCGTTTGGA 2460

ACCAGATATA AAGAGCAAAT ATTATAAAGA AGCAAAAGAT GTATTAAAAA ATATTTTAGA 2520

TTTGTAACAT TTTTTTATAT TCCCCCGGGG GAATATAAAA AAGGGAGTTT AAAATGAAAA 2580

TAAAAGCCGA AAAAGATAAA GAAGCATTAT TTAGTAATCG TTTTGGGGAT TGCAATGAAG 2640

AAACAAATTT AAATGACGAT CAAGATAAAG AATTGGCAAA TTATAATAAT CTAAAAGAAC 2700

AGCTTAAGTA TAATTTAAAA GATGATATTA ATAATAAAAT TCAAAGAATG AAAATATTAT 2760

ATGAAATTAA ACAAAAAGAA TTATATAAGT ACGATGGTTT TGCTCGTTTT AATGATTTTA 2820

TAAAATCTTT TGAAGTTGCA AAAAGTCAGG CTTATAGGTA TTTAAAAATT TATCAAAAAG 2880

TTCTAGAGGG TAAAGTGTCC ATTGATAAAA TAAAAGAAGT GGGCTTTAAG GCTATATTAA 2940

GAGATATAAA GGCCAAAGAT TCTTTAAACG AAGATAACCA TAGTGAATCT GAAGGCGCTA 3000

ATGAAAGCAT TCCTATTAGA ATTTTAGTAA AGGATAAAGA ATTATATAAT TTCTGTAAAC 3060

AAGATACTAA AAGATTGTAT TTTATTATTG AAAAAATTTA TAAAGAAAAG AGAGATGTTT 3120

TATCTGAGCT TATAATTGAG TATGAAAAAA ATAAAAAATA AAAAATAAAA AATAAGCCTA 3180

TTGATTAATA ATGTATATTA TGATAGCATT TAATCAGGTT GGGATTTAAT TTCCTATACT 3240

ATTTGTTCGT TGTAAACGGA ACTGCCCATA TCGGGCTTTT TTGCTATATA TAATCTTAAT 3300

CAAAAGTAAC TATTAACTAT TAACTATTAA CTATTAACTA TTAACTATTA ACTATTAACT 3360

ATTAACTATT AACTATTAAC TATTAACTAT TAACTATTAG TAGATTTAAG TTTTTCCCCG 3420

ACTTAATTTG AA 3432

(2) INFORMATION FOR SEQ ID NO: 35:

(i) SEQUENCE CHARACTERISTICS:
(A) LENGTH: 3398 base pairs
(B) TYPE: nucleic acid
(C) STRANDEDNESS: double
(D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 35:

ATAATAATAA TGATTATGAA ATGCTCAAAA ATCTTGGTAA AGGGGTTTTA ATGAAAAATG 60

CCAATGAATT TCTTAAAATT AATTTAGCAA AGAATGAAAT AACAAGATTT AGTAATAATG 120

AGGATGGCGT TGCTAGGTTT TTAATTGATT TTTTTAAGCT TAATATTAAA TATTAATAAT 180

TTGTATTTAA ATGTTTAATC CATTTGATTT ATTTTTAGCA GGATTTTCTA TTTAAAATAT 240

AAATTTTTTA CTTATAATGT ATTTTTGAAA ATTTATTTAT TAAAATATTG GAATAAGTAT 300

TGACATGGAT TAAACAAAGA TATATATTAT TTTATGTTGy ATAAACAAAT TGGCAAAATA 360

GAGATGGAAG ATAAAAATAT GGTCAAAGTA ATAAGAGTCT ATGGTGAATG CCTAGGAGCT 420

TTAAGGCGAA GAAGGTCGTG GTAAGCTGCG AAAAGCTTGG GGGAGAAGCA AACATTTATT 480

GATCCCAAGA TTACCGAATG GAGTAATCCA GCTAGCAAGA TGCTAGCTAT CTATTATTTA 540

AATAATAGAG GCGATACCAG GGGAAGTGAA CCATCTAAGT ACCCTGAGGA AAAGAAATCA 600

AAGAGATTCC CTTAGTAGTG GCGAGCGAAA AGGGAGTAGC CCAAACTTTA AATGTGTCAA 660

GCTGCAGAGC GTTGCATTTA TGGGGTTGTA GGACGTTTAG GCTTAGTCTG TAATAAGCAA 720

AAAAGTTACA AAATATTTAT ATAGAAGAAT AATCTGGAAA GTTTAACCAA AGAAGGTGAT 780

AGTCCTGTAA TTTAAATGTA AATATCTTTT TAAAATGTTC CTGAGTAGGA CGAGGCACGA 840

GAAACCTTGT TTGAAGCTGG GGAGACCACT CTCCAAGGCT AAATACTAGA AAGCTACCGA 900

TAGAGAAGAG TACCGTGAGG GAAAGGTGAA AAGAACCCCG GGAgGGGAGT GAAATAGAAC 960

TGAAACCGTA GACTTACAAG CAGTCAAAGC CGTAATTTAT TGCGGTGATG GCGTGCCTTT 1020

TGCATAATGA ACCTGCGAGT TATCATGTCT AGCAAGATTA AAGCATAGAA GTGCTGGAGT 1080

CGAAGCGAAA GCGAGTCTTA AAAGGGCGAT TTAGTTAGAT GTGGTAGACC CGAAGCCGAG 1140

TGATCTATTT ATGGCCAGGC TGAAGCTTGG GTAAAACCAA GTGGAGGGCC GAACTCTAGT 1200

CTGTTTAAAA AGGCAGGGAT GAGCTGTGAA TAGGAGTGAA AGGCTAAACA AACTCGGAGA 1260

TAGCTGGTTC TCCCCGAAAT GGATTTAAGT TCAGCCTTAT TTTAGTTTAA TAGAGGTAGA 1320

GCACTAATTG AGCTAGGGCC TGTCAAAGGG TACCAAACTC AGTTAAACTC CGAATGCTAT 1380

TAAATGATGA ATAGGAGTGA GACTATGGGC GATAAGGTTC ATAGTCGAGA GGGAAACAAC 1440

CCAGACCAAC AGCTAAGGTC TCAAAAATGT GTTAAGTGGA AAAGGAGGTT TAGGTACGTA 1500

AACAGCCAGG AGGTTGGCTT AGAAGCAGCC ATACCTTTAA AGAGTGCGTA ATAGCTCACT 1560

GGTCGAGTAC TTAAGCGCCG ATAATGTAAC GGGGCTAAAC ACATTACCGA AGCTTTGGAT 1620

CTTAACGAAA GTTAAGATGG TAGGGGAGCG TTCTGTAAGC CAGAGAAGTT AAGCTGGAAA 1680

GTTTGATGGA GGTATCAGAA GTGAGAATGC AGGTATGAGT AACGAAAAAA TGGGTGAGAT 1740

TCCCATTCGC CGAAAACCTA AGGTTTCCTG GGTAAAGGTC GTCTTCCCAG GGTTAGTCGG 1800

CCCCTAAGGC AAAGCTGAAA AGTGTAGTCG ATGGGAAACG GGTTAATATT CCCGTACCTC 1860

TTATAGTTTC GATGGAGTGA CGCATGAGGT TAACTACTGC TAGGCGATGG TTGTCCTAGT 1920

TTAAGCATTA AGGCGATGAT CTTAATAGGA AAATCCGTTA AGAGAGCTAA GATGTGATGA 1980

TGAGTGCTAT TTAGGTAGCA TGAAATGTAG GTAGTCAAGG TGCCAAGAAA TAGCTTCTAA 2040

GGTTAGGCTA TAAGGGACCG TACCGCAAAC CGACACAGGT AGGTGGGATG AAAATTCTAA 2100

GGCGCGCGAG AGAATCCACG TTAAGGAACT CTGCAAAATA CGTACGTAAC TTCGGGATAA 2160

GTACGACCTA AGCAATTAGG TAGCATAAAA ATGGTCCAAA CGACTGTTTA CCAAAAACAC 2220

AGGTCTCTGC AAATCTGTAA AGAGAAGTAT AGGGACTGAC ACCTGCCCGG TGCTGGAAGG 2280

TTAAGAGGAG ATGTTAGTTT ATGCGAAGCA TTGAATTTAA GCCCCAGTAA ACGGCGGCCG 2340

TAACTATAAC GGTCCTAAGG TAGCGAAATT CCTTGTCGGG TAAGTTCCGA CCCGCACGAA 2400

TGGTGTAACG ATTTGGACGC TGTCTCAACG TGGAGCTCGG TGAAATTGAA GTATCGGTGA 2460

AGATGCCGAT TACTTGTGGT TAGACGGAAA GACCCCGTGA ACCTTTACTA TAGCTTGGTA 2520

TTGAGATTTG ATTAAATATG TGTAGGATAG GTGGGAGACT TTGAAGCTAT CTCGTCAGGG 2580

GTAGTGGAGT CAATCTTGAA ATACCACCCT TGTTTAATTA GGTTTCTAAC TTATAGAAAT 2640

ATGAGGAGAG TGCCAGGTGG GTAGTTTGAC TGGGGCGGTC GCCTCCTAAA GAGTAACGGA 2700

GGTGCGCAAA GGTTACCTTA GAGTGGTTGG AAATCACTCT GTAAGTGTAA AGGCATAAGG 2760

TAGCTTAACT GTAAGACTGA CAAGTCGAAC AGATACGAAA GTAGGTCTTA GTGATCTGGC 2820

GGTGGCAAGT GGAAGCGCCG TCACTTAACG AATAAAAGGT ACTCCGGGGA TAACAGGCTT 2880

ATCCTTCCCA AGAGTTCACA TCGACGGAAG GGTTTGGCAC CTCGATGTCG GCTCATCGCA 2940

TCCTAGGGCT GGAGCAGGTC CTAAGGGTAT GGCTGTTCGC CATTTAAAGC GGTACGCGAG 3000

CTGGGTTCAG AACGTCGTGA GACAGTTTGG TCCCTATCTG CCACAAGCGT TGGATATTTG 3060

AGAGGAGCTA TCTTTAGTAC GAGAGGACCG AGATGGACGA ACCTCTAGTG TrCCAGTTAT 3120

CCTGCCAAGG GTAAGTGCTG GGTAGCTACG TTCGGAAAGG ATAACCGCTG AAAGCATCTA 3180

AGTGGGAAGC CTTCCTCAAG ATGAGATATC CTTTAAGGGT CCTGGAAGAA TACCAGGTTG 3240

ATAGGTTAGA AGTGTAAGTA TAGCAATATA TTAAGCTGAC TAATACTAAT TACCCGTATC 3300

TTTGGCCATA TTTTTGTCTT CCTTGTAAAA ACCCTGGTGG TTAAAGAAAA GAGGAAACAC 3360

CTGTTATCAT TCCGAACACA GAAGTTAAGC TCTTATTC 3398

(2) INFORMATION FOR SEQ ID NO: 36:

(i) SEQUENCE CHARACTERISTICS:
(A) LENGTH: 3203 base pairs
(B) TYPE: nucleic acid
(C) STRANDEDNESS: double
(D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 36:

GAGAAACCAA GAATTAAATG TAAAAATAAA GATCGTTTCA TAAAGATTGA AAAAGAAAAT 60

GACAAAACAA TGTATCATAC AAAAATAATG ATGGATATTT ATAAATTAGG AATTGACAAT 120

AAAAGAAATG AATGTCGTAT ATCATTAAGA ACACTATTTA ATCAAATGAA AGTAGAAGAA 180

GTTCGTTTAT ATTCTATAAA AGAAGGGGAC AAATTTTTAG GTATTTACTA TGGATATAGA 240

AAACCTATAA AAAACATTTT CGTAAAATAT GAAATAAACG GAACCATAAA GTCATATGGA 300

TTATCAAAAG CACATTACAT AGAATTTAGA TTTAAAAAAG GAAGCGTTTT TTGTTACTTT 360

AAAGGATTAT TTCGCTTATT AAAAAAAGAA AAAGAAAATA CACCATATAA TATGGCTTGT 420

ATTGATATGT TTACAAAACT AGAGAAACAC GTATATGAAT TTTACGGTAA AAAATATCCA 480

GAAAAAGGAA TAATTATAAG ATGGATAGAA AAAAATCAAA AATAATAACA ATTGCAAGCC 540

TTAAAGGGGG CGTTGGTAAA AGCACAACTT CAATAATACT TGCAAATCTA TTATCGAAAA 600

AGCATAAAGT ACTTTTGATC GATACAGATG ATCAAGCTGC TACTACAAGC TATTATTATA 660

ATGAATTAGA AACAAAAAAT TTCGATATAT CTAAAATGAA CATAGGAAAT GTTATAAAAG 720

ACGGTACAGA TATTAATAAA AGCATTATTA ATGTTGAAAA TAACATAGCT TTGATACCCA 780

GTTATATAAC AGTCGATGAA TTAAATGGAG AGTATTATTA TGATAACCGG CATCTTCCAA 840

TTGAATTTTC ATTAAAGACG AAATTAAATT CCATAGCAGA CAACTATGAT TATATTATAA 900

TTGATACTAA TCCCAAAAGG AATTTCACAT TAAAGCTTTC CCTAATTAGC AGTAATTATG 960

TAATATCTCC AATGACGGCA GAAAAATGGG CAGTTGAAGG ATTTGAAACA TTAAGAAGGT 1020

ATATAAAAGA AGTTGCTGGA ATACCAATAT TTATTGTTAT TACAAGGTTT AAAAAAAATG 1080

TTACCCACAA GCAATTAATG GAAATAGTAA GCATGAAAAA CGGGTTTTTG GGATACATAA 1140

GCGAAAGAGA AGATTTAAAT AAAAGAATAG GGTGTAATGA AAAATTTGAT TTTTCAAAAG 1200

ATTACATTAT TGAATATAAA AAAATATTGG ATGTTTTTTT GGGAAAATTG TAAGAATTGA 1260

CAAACTTAAT AAGTCCGGCA TGCCGGACTT ATTGGAAATA AGGGCAAAAA TATGAATAAA 1320

AAAAACATTA ATTTAAAAAT TAATAAAAGA ATTTCAGAAA ATAATTTAAA TTATATTCTT 1380

GATCAAAGCA ATGAGAATCA AAGAAAAGAA GAATTTGAGC GATTAATTAC ACAATTAAAA 1440

AATAATATTA AATCAGAAAT ATACAATATT ATTGATACCA TGAAGATCCT TAAGAAAATA 1500

AATGACAAGA GGCTCTATTT AGAAGGAGGA TATAAATCTT TTAAAGATTT TTTATCAGAT 1560

TTTAAATTAG CAAAGACACA GTCTTATGAA TATATAAAAT TAGCCGCTGC AATTGAGGCG 1620

GGAATATTAG AAAGAAAATT TTATTACCAA TAATGGAATA AGGGCCTCTA TAAGATATAT 1680

TAAAAATCAA GCAAATGGTA CAATAAAAAA ATCAAAACAA AATCCAATAA AACCATTAAG 1740

ATTTCAACTC AAGAACCAGG AAAGTTATGA CTTTTATAAA AGCAATTCTA GGTTTGTAAG 1800

TTTTATGATG GATGAGATTT TTAAAAATCA AAAAGATTTT CTTAATAAAC TTTTAAAAAG 1860

ATATAAGGAA TCAAAGGGAC AATAAGAAAA TTTTATAAGC AATTTAATCT TTAATATTAT 1920

TGAAATATAA AATATAAAGT TAGAAATTGT AAATAATTGA TTTAACAAAT AAGGAAATAT 1980

ATACAAAAAA GCAACTGAAA ATTTAAAAGA TCATTTGCTA AGCAGAGGAA TTTTATTTGG 2040

TTAAAGTATT TGAAAAAAAA TTAAATATTA TCAAAGAAAA AGGTAAATTG ATCTCAATTG 2100

ACAATAAATT GTCAGTAATA AGCAACGAAG AATGTTTTTA ATTTTGATTA TAATAAATTT 2160

TGCAAAGAAG CACGCTTAAT AAGCACGGTG TAATAAGTTG GGCAATATCT TGATTAGCAA 2220

CAATGTGTTT TGATAATCAA TGGTAAGACG AACACTTAAA AATATTTTAG AAGGAACAGG 2280

GCATTTAAAG TAAATTTATA TATATTTAAG AGTAAGGATT TGAATAATTT ATGATTTAGA 2340

GCTGTAGAAT TTATTAAAAT ACAAGAGCAA CTATTAAAAA AATGGGCAAA TTAGAAACTT 2400

TAGCAGGGGT AGTGATTTAT AAGGACGGGG TGTTAATGAC GATATTATAT AGGCTCAAGG 2460

GCCAGCAAGG ACTACTATTG ATTATATAAA CACTGTTGTT TAAGATTTTA AAAAAAAGAG 2520

GCTATTTTAA AATAACAAAT ATTTTATAGA ATTTATATTC AAGCATAAAT TTAGGAATTA 2580

AAATCAATGA GTTTACTATG AATCTTGAAT TTGTTTTGTT AAATTCAAAA CCACCACACC 2640

CCTATTTTTT AGCTATCTAA TTAAGGGATC CATATGTGTC CCCTTTATTT TTAAATAAAA 2700

GATATATATT TAAAGACAGT TAGGCCTCTT TTAGGCATAT TTTTGTTTAA TAAAAAATAT 2760

TAAATTAGGG TTTATAATTT TTATAGATGA AAATAAAATA GAAGAATCTA ATTTAACTAA 2820

ACAATTTTTG TTTAGTTAAA ATGATATAGG GCTTTGCAAA GTAGATATAA TTAAAGAAAA 2880

TCTAAAATCG CTAAATAAAA CTATTTAAAC TAAGCCCCAT AATGAAAAAG TTTTAGTAAA 2940

AATATTAAAG AATATTTTTA CTAAAATAAA AATTAAACCA GCATTAATAA TACTTACATT 3000

AGATGATTAG CTACTTTTTT TAATTAATAA ATTTTGCATT TAAAGTTCTA TTCCACTTAT 3060

AAATATTGAC TATATCAATA ATTTTTCAAG CATTGGTACA TTTTATATTC TAAATATTTC 3120

GTTTTGTCGC TAATTTGTTG ACATAGGAAT TATAAAAAGG CCATCATCTT TTAAATTAAA 3180

AAGTAAAATA ATACTAATAA ATA 3203

(2) INFORMATION FOR SEQ ID NO: 37:

(i) SEQUENCE CHARACTERISTICS:
(A) LENGTH: 3189 base pairs
(B) TYPE: nucleic acid
(C) STRANDEDNESS: double
(D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 37:

TTGCTAAAAA TATATTTTTT TATATCTtGG CTCTACTACA ACTCCTTTAT TTTTAGAGTA 60

ATAAATAAAT CTGCTGCCAA ATAACCTATA AGATATTTTT TAAACATAAC TTGGATCGAT 120

ATTTATAGAT TTAAAAATTT TTAAATCTAT AATAACTTCC AAAATACAGA ATCTTCACCT 180

AGATTATAAT TGTTTATAGC ATATTCTTCT TTTGAATATG TAAATTCTTA AGCTTATTTA 240

AAAGCTTTTT TCTTGTATAG TCAATTAGCT ATTTTAGCTT TTCTGGTTTG AACGCCAATA 300

CTAAAATTAT TGAAAATATC TTTTAAAGAA ATCCCATTGT CATAAATATC TTTATTTGAA 360

AAATCTTTTT AGTAAAAAAA TAATGGGGAT ATTTGTATTC AAGTTTTTCA AAATTAATAG 420

TAAAGATATT GTTTTTTCCT AAAAACCCAT ATTTTTTTTC TCTTTTGCCT TTAATACTTT 480

TGTAAAATAC TTTAATCAGC TTACTTTTTT TAGTCCATCT TATATTTAGT AAAAATAGCA 540

ATAGTAACTC CGGTTTGGAT ACAAAATACA TTCTCACCTA TACTGCCATC ATCAGTTTTT 600

TTCTTTCTTG AACTATCGTG TAAATTTAAT ATATAAATTT CATCAAATGT TTTTAAGAGG 660

TAATCTCATA CCTCTGAATG CCACCTTGTC AAGGTATCCA TTTGTTTGTT ATGATACCTA 720

GCAATCCTtC GTTAGAACTT TTAATTCTAT GTTCTGCAAA TCTAATAAAC TTAGTGCGGT 780

CATCATTAAG TGGCCTTAAA TTTTTTTCAT TTATAAACTT ATTTTCTATT TTTTTATAAT 840

CATTAACTAA ATTTAATATA TATTCATTAT TATTTTTTGA ATCTGAATTA TAAGGAGGAT 900

TTCCCAGTAT GACAAGTATT TGCTTTTCTT TAGCTTTATT TGTAAGTTTA TTTTCTTCTC 960

CAATTGCAGG GAAAATACCT TAAAAGATTT TTGATCTGAA GGGTCTGCTT TATCAATAAA 1020

ATTAGTCAAA AATATTTATA ACTTTATGTT TTCATTATTT AAGCTGTCGC TACAAATTTT 1080

TCTTTCAAAT ATTGACTTGG CTTTAAATGA TCAACTACAT AAGGAACCAT TAAACATTCA 1140

AAACCATAGA CATTTTTAAG TATGTGAAAA TTAATGTAAT CTTCTTGTTT TTCAGAGTCT 1200

ATTGGAATTT CGTTTAATAT AATTTTAATT ACTTCAGGTA AAAATGTGCC AATACTTATT 1260

ACAAAATCAA ACACTGCAAC TTTATCCCTA TTTTTAAAGC CATAATTTAA TTTAATTTTT 1320

TTAAGTGCCT TATGTAAACT GTTAACAATA AAACTTACAC CCGAATAGGA AGTGTACTAC 1380

ACTCCTTTGG CCTTTCTTAA TTTAACATCG TACTTAGCTA GAAAATCCTC ATAAAAATAA 1440

AGATAGGGGA TCTTTTGAGC TTGTTTTGGC TCTTGTAAAA GAAAATTCTT TAAAAAATAA 1500

GCTCTGTATC AATTTATTTA TTACATTAAT AATTTCTTCT AAAATCTATC TTGGACTATT 1560

ATATTCATTA TTTGTATCAA TATCACTAAT TAATTTTAAT ATATCTCTTA TAAGCGAAAA 1620

GTTAGAAAGT ATAAACTTTT TAATATTATA AAAAGCTATT TTTCATTAAA TTTAAGCAAA 1680

GACATAATTA CCTTATCTTA AACTTTTAGA TAATTATAAT TGATTTTTAA GGAAATCAAA 1740

ATGATGCCCC AAAAGCTTTT AATAATAAAA AATTGTTATT CTTGCCAAAA GCTACTTAAG 1800

AAAAATAGTA AAATTTGTTG TGTTGTTTAT AGAACTAGAA ATAAATACCC CAAAACCCTT 1860

ATAACGAGCT AATAATTGTT TTTATTGGGA GGTTATTTAG GGATTGTAAA TTTTTATATA 1920

CAAAGCCTAA AACAAATTTT ACATTTTTAT ACTCATATCC TTTTCTATTT CTTGCTTGCT 1980

GTTTCTTTAT AATAACGAGT TTTTAATCAA TTAAAATAAA ATTTTTCTTT CTAACATCTT 2040

TCTTTCTTCT TTCAAACCAT CTTCmAGACA CTTTAATTTA TCTTTATAAC TCTTCtTTAT 2100

CTTTTGGTTA TTCtTCtTTT TTAAATTTcT CTTACTTTTT GTATTcTTTT TCAATCTTTT 2160

CTAATTCTTG TATtCCTTTT TTATTAACTT CTAATCTTAG ATTTTCAACA ATATTTTAAG 2220

CTACCAGATT TTTAGATTTC TCTTGTATCT CTGCTTGGCA TCTTAAATTA ATTTCTCTTT 2280

TATCTCTTCT TTTGTATCTA TAAAATCTAA AAATTCTTTA GCTTGTTTTT TTAAATCTTA 2340

TTCTAAATTT TGTATATCTA CAACTTATAT AATGTTTATA AGAACTTATT AATATAAAAA 2400

TAGCAGACAT AGTAAACTTT TCCTCATAAT TATCCCACCA AGCTTTAGGA ATATATTAAA 2460

TACCTATATT TACATATATA AAGCTTAAAA GCAATTTAAT TACATTGCAC ACTATAAAGC 2520

AAAAAAATAA AGCCCAAAAA AACTACTCCC CCCAGAGCAC AAATTCCAAA ATCACCAAAG 2580

CAGATCACAA TAAAGACAAT TAAATGCAAA CAATAAAAAT ACAAGATATT CCAACTTTAT 2640

TTAATAAAGT TGGAATAATT TTTTGCAACA TAAATTTTGA AAGCATTATC AAAATAAACA 2700

TTTATTAATA AAATTGTAAA AAAGAAAAAC CGATATTAAA TACAAAGCTC AAATATCTAT 2760

AAGTTAATTT AGTTTTACAA CAACTAGCAA TAAAACTACA TTAATAAAAT TAATAGATAC 2820

ATTAAATTTC AAATCTTCAT TACAGACACT ATAAAACAAA ATTTTGTATT TTACTTTTTC 2880

TTATTATTAT ATTAGTTACA ATATCAAGCA AAACATCTTT TCTATTCTAA AAAGCTCCTT 2940

AAAAATATAA TTTACTATTA TTATCTCTTT TTTACAAACT CACATAATTC TTTATCTCTC 3000

ATGAAAATTC TAATAGAAAT ATTCTTATTA TTTAATTTAC TTATATGATT ATTAAAATTT 3060

AATGATTTTT TTTCTAAACA TGTGTTTTTT TATATTTTTA AATCCTAATT CCTTAATCTT 3120

ATCAATAGAT AAAACTATCC TCCAAAACTT TTGACTAAAG TTTTAAATAA AAATAAGTTT 3180

GCnTTTAGC 3189

(2) INFORMATION FOR SEQ ID NO: 38:

(i) SEQUENCE CHARACTERISTICS:
(A) LENGTH: 3130 base pairs
(B) TYPE: nucleic acid
(C) STRANDEDNESS: double
(D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 38:

TTGTAAAAAG TTTTGTTTGA TATAAGTTCA CTTGTGCCCT TTACTATTTT TGTAGTTGCT 60

TTTACTTTCA GCTTCATTAG CAGTTGCAAG TTTTTTCATA CATTCATAGT AAAGCTCCAT 120

TTCTCTTATT GAACACTCCT TTATATATTC ATCAAGCTCG CTTTTTAAAG AATTAATTTC 180

TCCATTAACA ACTTGCTTAT TTTTTTTACT ACTTGCTTTA TTTAAAGCGT CAATTTCAGC 240

TCTTAAATTT TCTATTTTAG TATGCATACT AAAAAGCTCA ACACTAGAAT ATTGCTTAAA 300

TGCATTTATA AATCCTAATT CTAAATTAGC CCGCTCTAAA TCCAATTCGC TTCTAACTTT 360

CCTAGCGTTA ACTTCTGATT TAAAAGTTTG CGACAAAAGG TGTTCAAAAG TGTCTTCACT 420

AATTGATAAT CTAGAGTCAT CGCTAACAGA ATTTTCCCCA CTTTCCCATT TTTTCCTCAT 480

CCTCCACACA TTTACCCTAG AAACCCCCAA TTTATCCGCT ATTTCCCTAT CATCTAACAA 540

TCCTTCTCTA AAATATGCAA CATAATCATC AAAAGACCTT TTAACTTTTT TCAAAAAATC 600

CTCTAAAATA ACAAAATTAA CAAATTGTTG CTCTAAATAG TAAAGCAATT TATCAATTGT 660

TAACATTAAC TATTGTCTTG TTGATATCTA TTGACCACAG ATCTATCTTT ACAATTCTTA 720

TTAAACATGA ACCAGTATCA TTATTGTCGC CATTAAGAAG ACCCCTCATA AATTCGTTAG 780

CATTTAAAGA AAATAAAGCA AATATAAAAT AATATAAATT CTTTTTATTT ATCATGATTC 840

TCCAATATTA ATAAAATAAC AAGACTAGTA GCTAATCTTG TTATTCATAA TTTATGCTTA 900

TAAAAACCAT TTATTTTATT TCGAAATCTT TTTTAGCTTT TCTTAATAAA TATCTTATAA 960

GATTCTTTTT TCAAATTAAA ATCTAATCTT TGGGCAAATC AGCCAAAATT TGTTTTAAAA 1020

TTTGTTTAAC TGTATTTGCT TTATCTTCAG AATAATCTTT TTTAAAATTA TTTCTGGCGT 1080

TATCTCCATA TTTCTCAGCA TAATCAATTT TATCCGAATT TAATTGTATC AAATAATTAA 1140

AAATCGAATC TGGATAACTC CCTATAAGTC TAATCATATC CTCAGACAGG AAAATACTAT 1200

CAGTACTTAT CTTAATTTTT ATAAGATATT CAATAGCCTC AAGAGCGTCC AAAAAAACAC 1260

TTTTTTCTTT AATTCCAATT TTTCTTAAAT CTCCTCTAAT TCTAGGAGCA TCGGAAAAAA 1320

CATGACTTTT TTCATACTCA TTTTTAAAAT CATAATTATC TAGTCTTTTA TTTATTAGGT 1380

TAAAATCTTC TTTAGAAAAA GCTCTTTTAG TTTCTGTATA ATTTTCTTCT ATATTTGCAC 1440

TTAAACTTAC TACAAATAAA AACAAAAATA TTAACAGACT AATTTTTTTC ATATCCCCTC 1500

CTAGCTTTAT TGCCTAAATT TCAGCAATGT AAATGCTAAT AAACAATAAG ACTGATTGTT 1560

AGTCTTGTTG TTTATAATTT TTACTATCAA AACCCATTTT TTATTATTTT TTATCTTCTA 1620

TATTTTGAGG CTCTGCTAGC TTTTCAAGTT CTTCCTCAAT ATTTTTAAGA GCATCATCTA 1680

TAACCTTTTT TACAAAATCA TTAGTATTAG TACCATCATT AACAGAATAA CTTCCATTTA 1740

CACCCAATTC TTTAGCATAC TTTAAAGCTT TTTGTCCAAT ATTTCCTTGT TTTTTAACTT 1800

TCTCAGTACT TTCTCCAGTT GCAGATTCAA CTTGTTCTTT AAATTCTTGA AATTTCTTTC 1860

TAGCCTCTTC TAATTCTTTT TTTCTTTTAT CTATTTTATC CTTTAATTCT TGAATTTCTT 1920

TTTCTTTCTC TTCTTTTCCT TCTTGTTGAC CACCATCTTT TGCTTGCACC GCTTTTAATA 1980

CGGGTGTGTT ATCGTGGGAA CTTGCCGGCA ATACTGGTGG TGGATTAAAC AGACTGTTAT 2040

TAGGATCGTC ACCTTGCATT AACTCTTTAT CTAAAAATCC TTCAACTTTT TCTTTTACAT 2100

TTTGTTTTAA ATCTTCACTA CTCGCATCAA TCTTGCAAGA AATTATCAAA ACAAAAACTG 2160

CACAAATAAT TAAATTTTTC ATTTTCTTAT TCATAAGTTA CTCCATAAAG TACTAATATT 2220

ACCACAACAC CAAATAATTG CAATATTTCA AAGATTTAAA TATATAATTT TGTTACATTC 2280

AGCTGTTACA TTTTAACAAA ACACAAATGT AATTTTAACC AACTCGCCAA AATCTCTCCA 2340

TTGCAAATGC TCTACTCATT ACAAAAGATT ATAAAATACA TACAAATTAA ATTTTCAAGT 2400

CTTTGCTATA TATTACACAA AGTATACTAT CTTTCTTGTG TACCACCCTC AAAAATCACT 2460

ACTTCTGTTT ATTACACCCA CTCTACAGCC CAGATTTTGC ATGCAATGAG AACACTCCAA 2520

ATTTGACTAA AATTTTTCGT TTTTAGTAAA ATATAATTTA CATTTTTTAT CTATTTTTAT 2580

TACTTTTACT TAATTTAAAA GTAACAACTT CAAGGAGAGG ATTTTATGGA CAATAATAAT 2640

TCTTTTAATT TAAATAATTT CAATATGGAT TTTACGCTCA AACTATTTCA AGAATACCAA 2700

AAACTAATAA ATGAAAACAA AATTCTTAAA AATTCACTAA AAAATTCATC TAAAAGTAAA 2760

AAAGAAAATT CAAAACCAAC TCCTAAGTTT TATTTAACCC CTAAAAGTAT TAAATTAATT 2820

CTAAAATGTG CCAAAACCTT AAAACAAATT GACCCAATTT CTGGTTGGTT TGTGCATCTA 2880

CTCTTAATAA GTGGATGTAG AGGCACTGAA ATGCAAAAAG TAAAAATGCA AGATATTTCA 2940

ACTTTTTTAA GCAAAACCGG AAAAACTTTA TATACTATTA AAGTAAATGT GGCAAAAAAA 3000

AGAAATACCT CTTGTATTAG AGAAATTGTC ATCAACTCAG AAGAGTTCGA GGCTATCCAA 3060

ACAGCACATA AAAATCATTT CCAAGAAAAA ACTCTTGACT CAAGGCGTAC TTATCTTTTC 3120

CAAAAGAGCA 3130

(2) INFORMATION FOR SEQ ID NO: 39:

(i) SEQUENCE CHARACTERISTICS:
(A) LENGTH: 3029 base pairs
(B) TYPE: nucleic acid
(C) STRANDEDNESS: double
(D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 39:

AGTTTTCCCA TTTCGCTCCG TAACTTGTGA TTACACTATC ATTTGmGwcy CCTGATATkT 60

cTGTGGAkaT TATTGTTgAA GmTTTTCCtC TTTTcCTCCT TATTTTGTAa TAAaTAtTAT 120

ATgCAAAAAC TATTTTTGCC AACTTTTTTA CAAAAAATTT TACAAAAAAT TAGTTGGGCT 180

TATTTAAATT CTCTTGTTAA AGAACTTAGC TAAACCCAAC TTAATTGGAT TTTTTTGATA 240

GCAATATATT TTTTTAGTTA TACATATACA TATGTATAAC TAAAACTATT TTTGTCAGGC 300

TTTTTACAGA AATTATTATA ATAAATAAAA GCTTTATTAA ATTCTCATGT TAAAGAGCTT 360

AATAAAGCCG CGGGCTTAGC TTAATTTATT TTAGATAAAC TCACTAAACA ATTAAATTCA 420

GTTCAATTTA GAATTAAAAT TATTTTGTAT TTATTAAAAT AAAAGAACCT ATTTAAATTA 480

TCTTGTTAAA AAATTCAAAT AAGTTCTACT TTAGAGCTAT ATACTAACTT ATTACTTTAT 540

AAAATTTTAA TCATTTTCAA TTGAAAAAAC ACTTATTAAA TATAGAATAG ATAATTGGGG 600

CAAACGTTAT TCCCATTATT AGAATTACTT GTATTGTTCT ATTGCTTGCA TTAAGTTCAT 660

TTTTTTAAAT ATCTATTTTA TTGTCTAGAC TAGATATATC TTTTTGCAAA GTTTTTTCTA 720

CACTATCTAT TTTAGTATTT ACACTATCTA TATCTTTTTG AAAATTCTTT TCTACATTAA 780

TTATTTGTTG TTCTAAAGAA TTCATTTTCT CCTTTAAAAC TTCAAAGTTG TAATTGTCGT 840

TCTGAAGTAA AACAAAATCT ATTGCCTCTT CACTAAACCC CTTATTTAAA AATTCTAACC 900

TTATATTTTC TATTTTAAGA GCATTGTAGG CTAAATTACT CATAAAATCC CCTTTATTAT 960

CCTTTTAATT CTTTATATTT TTTTAAAAGT TTATTAATCA AATCTTTTTG ATTTTCAAAA 1020

ATCTCGTCCA TCATAAAACT AGTAAATTTG GCATTTTTTT TATAAAAATT ATAACTTTCC 1080

TGTTTTTTAA GTTGAAATCT TAGGGGTTTT ATTGGATTTT GCTTTGACTT ATCCTCTTTA 1140

ATTTCTACGT TTAATATATT TCTATATACA CCCTTTAAGC CTTTTTCTTT AATATCATTT 1200

ATTGATATGC TCCCCTCTAA TACTTTTCTA TAAATTTTAA GGTATAAAAA AGCCTGACTT 1260

CTAGATATTA TAAATTCTGA CAAAAAGTCT TCAAATTTTT TATAACCATC AATCAAATAA 1320

AGTTTTTTTT CTCTTATTTT ATATAGGATT TTCATTGTTT TAATTTTATT CTCAACATCA 1380

TCAACAGTAA TTCTACGAAG TTGATCCTTA TAGCTTTTAT ATTCAAGTTC CTCATTTTCA 1440

AATTCTTGAA CATCCTCAAT TCTATTATTT AATAATATTT TTTTTACTTT TAACTTTGAC 1500

ACTTAATCCT CCTAAGTTTC TGATTTATTT TTAAAAGTCT TCCGGAAGAC TTTTAAAACA 1560

TATTGTTTAA TATTTTTTTT ATTTCTTGAT AATAAATTTC TTTATTATTA GGCTCTTTCA 1620

ATTCATTTAT AAAAACCTTA ATTGAATTAT AAAAATGAAC TCTTCCTTTA ATAAGATCTT 1680

TGTATTCTGA CTGCAAAATA CTTTCAATAT CTTTATACGT ATTTCTATTT TTTATAAATT 1740

GATTTTCTAT TATTAACACA TCAATATTTT TCTTTCTTAT TATTTCAACT TCCTTTATTT 1800

CATTCATTAA TATTGGCAAA GACTCTACAG ACCACCTTTC TGCTTGAATA GGTATTATAA 1860

CTTTATGTGT AATGTTTAAC GCATTAAACA ATAAAGAACT TAAACTAGGG GGAGTATCAA 1920

TTACTACATA ATCAAAATTA TAATAATGTA AATTTTTATC AAATATATGT TCTAACATAA 1980

GCTCTTTATA AGGAATATCT CCTTTTTCAA ATTTACATAA AATTGGATGG GCCGGAATAA 2040

TATACATATT ATTATTTATT GAATTTATAT ATTCATTAAA AGCAATGTTT TGATCTCTTT 2100

TTAAAAGATA ATAAACATTA TTCAATTCAA TATTTCTGAT ATATTGTAAA AAATAACTGG 2160

TTAAACTATT TTGAGGATCT AAATCTACAA TCAACACTTT ATTGTTCATT TCGCTTAAAA 2220

TATATGAAAA TATAATTGAC AACATGCTTT TGCCAACACC GCCCTTAATT GACGCTATTG 2280

TTATTATTTT AGGTTTTTTA TTATCCATTT TATTAACGGT CCTTGTTCTG GGTATTTTTT 2340

CCCATAAAAT TTATATACTT GTTGTTCTAA ATCTGTAAAC ATACTAAATA AAACTTTGTT 2400

GTAATGATTA TTTGTTCTTT TTTTATCTAA TAATCGATAT AATCCCTTGA AATAGCAAAA 2460

TACACTTCCG GCTTTAAATC TAAATTCCAT ATAATATGCC CTTGCTAATG CATATGCTTT 2520

TCTAGTTCCr TTTATTTkAT ACTTTATTAA wGGyTTTTTn ATTGGTTTTC TrTAGCCATA 2580

AAAAATACCA ATAAACTTAT CTCCTTCTTT AATTGGGTAy AArTGAGTTT CTTCAACAAT 2640

yCTTTCCCCA TTAAATAGGG CCCTTAATGA TAGTCTAAAT TCATGTTTTT TTTCATATAC 2700

TCCAAATTTG TAAATGTCCA TCATTATTTT TGTATGGTAC ATTGCTTTAC CATTTTCTTT 2760

TTCAATTAAA ATAAAGCGTT CTTTATTTTG ACATTCAACT TTACATTTGC CCTTTTTTAT 2820

AGTTTCAATA GGCTCCATTG CACTTTCCAT ATTTAAATCC TCATATAGCC TTTATGTTAA 2880

ATTCTTCTGT GGTTAAAGAA TTTTTTTGTT TTCTTATTAT TTCCAATAAT TCAAGATAAT 2940

ATGTACCAAA TACTTTATTG TATTCTATTT TCTTTTGGTT ATTCAAATAT TTTTTGGTAA 3000

TTGGCTTTAG AATTTCAATA TTTgTTTCC 3029

(2) INFORMATION FOR SEQ ID NO: 40:

(i) SEQUENCE CHARACTERISTICS:
(A) LENGTH: 3000 base pairs
(B) TYPE: nucleic acid
(C) STRANDEDNESS: double
(D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 40:

CnCACAGnAC CTTTTAAAAG AGTATTCTAT TTTCTTAATT TATAAAAAAA TAAAATCAAG 60

GTAAGAACTA TTTAAAATAA ATCTTGTGAA TCTTTAGTAG TGAATGATTG TCTGTCCTAG 120

TAACTTAGAA CTTAGAAAGT TAGCAAAGCA AACTTTCCAT CCTTCTTCAT CTCATTACCA 180

AAATCCGCAT CCTCATCACC CTTTCCAATA GCAGCAGCAA TCGGATTTGT AGCCTCCCCA 240

GGCTTCTCTC CATCCTGCTC AGCCTCACCA GCAGCCTTAA CAATCGCACT TAATATCTGC 300

TCCCCACTAA CAGCACCAGC CGCCTTGCTA GCAGCCTCAC TGTCCCCAGC ATTAGCATTA 360

TCAACTTTCC CAAACAACTT CCCTGCCTTT TTATTATTCT CCCCTGCAGC AGCAGCAACT 420

TTCAGCTTTT CACTCCCCCC AGCAGCTTCA ACAATCTCCT TTATTCCCTT AGCAATCCCC 480

GTCACACTCG CCTCATCAGC AGCCTTCGCA GCACTATTAT CAGCCACAAC TTCTCCAATT 540

GCATCAGTAC CACTTGAAGC CCCCTCAGCT GTCTTTACAC AGCTTTTACC AGCTTATCCA 600

ACAACTCGCT AACTTCTTTA ATAGCCCCCT CAGCCTTCTC TTTCTCACCA CCACCACTCT 660

TCACAGCAAA CTTTCCATCC TTAGCCATCC CCCTCAAAGC AATAGCAGCA GCAATCTGAT 720

CATCCTTCTT CATCTCATGA TTAAACTCCG CACCCCCATC TTTATCCCCA ATAGCAGCAG 780

CAATCGGATT TTTAGCATCC TCAGGCTTCT TTCCCTCCTG ATCAGCCGCA CCAGCAGCCT 840

TAACAATCGC ACTTAATATC TGCTCCCCAC TAACAGCACT AACAGCACCA GCCGCCTTGC 900

TAGCAGCCTC ACTGTCCCCA GCAGCAGCAC CAGCCTTCCC AAACAACTTC CCTGCCTTTT 960

CATTGCTCTC CCCTTTAGCA GCAGCAACAG CTTTCAGCTT TTCACTACTC CCCCCAGCAG 1020

CTTCAACAAT CTCCTTTATC CCCTTAGCAA TCCCCGTCAC ACTCGCCTTA TCAGCAGCCT 1080

TCGCAGCATT ATTATTATCC ACAACTTTTC CAATTGCATC AGTACCATTT GAAGCCCCCT 1140

CAGCTGTCTT TACAGTTGTT ACCAGCTTAT CCAACAACTC GCTAACTTCT TTAATAGCCC 1200

CCTCAGCCTT CCCTTTCTCA CCACCACTCT TCACAGCAAA CTTTCCATCC TTAGCCATCC 1260

CCCTCAAAGC AATAGCAGCA GCAATCTGAT CATCCTTCTT CATCCCCTCC TTATTAAACT 1320

CCGCACCATC ATCCGCATTA CCCTTCCCAA TAGCAGCAGC AATCGGATTT TTAGCCTCCT 1380

CAGGCTTCTT TCCCTCCTGC TCAGCCGCAC CAGCAGCTGC AGCCGTAACA ATCGCACTTA 1440

ATATCTGCTC CCCACTAACA GCACTAACAG CACCAGCCGC CTTGCTAGCA GCCTCACTGT 1500

CCCCATTAGC AGCATCACCA GCTTTCCCAA ACAACTTCCC TGCCTTTTTA TTATTCTCCC 1560

TTGTAGCAGC AGCAACTTTC AGCTTTTCAC TCCCCCCAGC AGCTTCAACA ATCTCCTTTA 1620

TCCCCTTAGC AATCCCCGTC ACACTCTCCT TATCAGCAAC CTTCGCAGCA GCATCATTAG 1680

CCACAACTTC TCCAATTGCA TCAGTACCAC TTGAAGCCCC CTCAGCTGTC TTTACACAGC 1740

TTTTACCAGC TTATCCAACA ACTCGCTAGC TCCCTTAATA GCCCCCTCAG CCTTCCCTTT 1800

CTCATCACCA CTCTTCACAG CAAACTTTCC ATCCTTAGCC ATCCCCCTCA AAGCAATAGC 1860

AGCAGCAATC TGATCATCCT TCTTCATCTC ATGATCAAAC TCCGCACCAT TCTCCGCATC 1920

ACCCTTCCCA ATAGCAGCAG CAATCGGATT TTTAGCATCC CCAGGCTTCT TTCCCTCCTG 1980

ATCACCAGCA GCCGCACCAG CAGCCTTAAC AATCGCACTT AATATCTGCT CCCCACTAAC 2040

AGCACCAGCC GCCTTGCTAG CAGCCTCACT GTCCCCAGCA TTACCAGCAT CAACTTTCCC 2100

AAACAACTTC CCTGCCTTTT TATTATTCTC CCCTGTAGCA GCAGCAACTT TCAGCTTTTT 2160

ACTCCCCCCA GCAGCTTCAA CAATCTCCTT TATCCCCTTA GCAATCCCCT TCACACTCTC 2220

CTTATCAGCA GCCTTCGCAG CAGCATCATC AGCCACAACT TCTCCAATTG CAGCAGTACC 2280

ACTTGAAGCC TCCTCAGCTG TCTTTACAGC TGTTACCAGC TTATCCAACA ACTCGCTAAC 2340

TTCCTTAATA GCCCCCTCAG CCTTCCCTTT CTCATTATTA TCCTTCTTCA CAGCAAACTT 2400

TCCATCCTTA GCCATCCCCC TCAAAGCAAT AGCAGCAGCA ATCTGATCAT CCTTCTTCAT 2460

CTCATCCTTA AACTCCGCAC CATTCTCATT ACCCTTCCCA ATAGCAGCAG CAATCGGATT 2520

TTTAGCCTCT GCAGGCTTCT TTCCCTCCTG CTCAGCCGCA CCAGCAGCCG TAACAATCGC 2580

ACTTAATATC TGCTCCCCAC TAACAGCACT AACAGCACCA GCCGCCTTGC TAGCAGCCTC 2640

ACTGTCCCCA GCATGAGCAG CATCACCAAC CTTCCCAAAC AACTTCCCTG CCTTTTCATT 2700

GCCCTCTTTA GCAGCAGCAA CTTTCAGCTT TTCACTCCCC CCAGCAGCTT CAACAATCTC 2760

CTTTATCCCC TTAGCAATCC CCTTCACACT ATCCTTATCA GCAGCCTTCG CAGCATCAGC 2820

CACAACTTCT CCAATTGCAT CAGTACCACT TGAAGCCCCC TCAGCTGTCT TTACAGCTGT 2880

TAcCAGCTTA TCCAACAACT CGCTAGCTCC CTTAATAGCC CCCTCAGCCT TCCCTTTCTC 2940

ATCATCATTC TTCACAGCAA mCTTtCCATC CTTAGCCATC CCCCTCAAAG CAATAGCAGC 3000

(2) INFORMATION FOR SEQ ID NO: 41:

(i) SEQUENCE CHARACTERISTICS:
(A) LENGTH: 2991 base pairs
(B) TYPE: nucleic acid
(C) STRANDEDNESS: double
(D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 41:

GAGAAAAAAC ATGAATTTAG ACTATCATTG AGGGCCTTAT TTAATGGAGA AAGAATTGTT 60

GAAGAAACTC ATTTGTACCC AATTAAAGAA GGAGATAAGT TTATTGGTAT TTTTTATGGC 120

TACAGAAAAC CAATAAAAAA GCCATTAATA AAGTATCAAA TAAACGGGGC TAGAAAAGCA 180

TATGCATTAG CAAGGGCATA TTATATGGAA TTTAGATTTA AAGCCGGAAG TGTTTTTTGC 240

TATTTTAAAG GGCTaTATCG ATTATTAGAT AAAAAAAGAA CAAATAATCA TTACAACAAA 300

GTTTTATTTA GTATGTTTAC GGATTTAGAA CAACAAGTAT ATAAATTTTA TGGGAAAAAA 360

TACCCGGAAC AAGGACCGTT AATAAAATGG ATACTAAAAA ACCTAAAATA ATAACAATAG 420

CGTCAATCAA GGGCGGTGTT GGGAAAAGTA CGAGTTCAAT AATATTTGCG ACATTATTAG 480

CTCAAAAATA TAAAGTATTA TTAATAGACC TAGATACTCA AGCATCTACT ACCAGTTATT 540

TTTGTAAAAA ACTTGAAAAT CAAAAAATTG ATCTTGTCAA TAAAAACATA TACAGAGTAT 600

TaAAAGATAC ATTAGATGTA AATAATGcAA TTGTAAATAT TaAAGAGaAT TTAGaTTTAA 660

TACCAAGTTA CATAACTTTG CATAAATTTT CAAATGAATT TATACCCCAT CAAGAGTTGA 720

GATTAAAAGA TAGTTTAATC TTTTTAAAGC AAGATTATGA TTATATAGTA GTAGATACTA 780

ATCCTAGTTT AGATTTTACT TTATCAAACG CTTTAATAAC TAGCAATTGT GTAATAGTTC 840

CAATGACGGC AGAAAAATGG GCAATAGAAA GTTTAGATTT ATTAGAATTT CATATTGAAA 900

ATTTAAAAAT AAAAATACCA ATTTTTCTTC TTGTGACAAG GTTTAAAAAA AACAATACTC 960

ATAAAGAATT ATTAAAATAT GTTGAATCTA GGGAAAGATT TTTGGGATTT ATTCATGAAA 1020

GAGAAGATTT AAACAAAAAA ATTGCGGGCA ATAATGAATT CAATATGGAT AAAGACTATA 1080

TTAATGAATA TAAAGAAGCA TTATCAAAAT TTTTTGAGAT ATATTAAAAA ATTTATTATA 1140

AAAAAAATCC AGATTCTGGA CTTTTTTGAA ATAAAGGAGA TTTTTTATGA AAATAGAATT 1200

AAATAAAAGA ATTTTGGCAT CAGGGATAGA TCCCGATGGT AAAAAAGAAG TGATTACCAA 1260

TGAAGATAGA ATTGCTCATT ATAATGCTTT GAAAGATAGA TTAAAGGCTA ATTTTAGAAA 1320

AGAAATATAT CATAAATTGG ATAGCATCAA AATTTTGAAA GAAATAAAGG ATAATCAATA 1380

TTATAAAATT GATGGATATA AAAAATTTGA CTATTTTATA AAAGATTATA AAATAGCTAG 1440

AAGTCAAGCT TATAATTACT TAAAATtTAc AACTGCGTTG CAAGAAGGAA TTCTTAAAGA 1500

AGATTATTTA ATAGAAAATG GCATTCATAA TTCTCTTGAT TTAATAAAGG ATAAAGAAAG 1560

TCCAACATTA AAAAAGTCTA AACAAAATCC AATAAAACCT CTAAGATTTC AACTTAAAAA 1620

TCAAGAAAGT TATGATTTTT ATAAAAGCAA TGCTAAATTT ACGGGATTCT TGTTAGATAA 1680

ATTATTTATG GATAAAAAAG AAATAATTAA AATAATTATG AAAGAATATA AACAATTAAA 1740

GGGATAATAT GGAGGTTGTA TGAACAATTT AGCTAACAGA ACGTTTAACA TAGGAAATAT 1800

AAAAAACGAA TTTTTAGAAA TAGGATTTAG CGAAGAGGcA ATAGATTTTG TTTTTCTTCA 1860

TAATGATAAT TATAACTTTG AGTTTTTAAA AGAGAAATTG ATTAATTTAG AGAAGAATTT 1920

ACAAAAAGAT ATATCTAATT TAGATATCAA AATAAATAAT GTTAAAAACG AACTTAATGC 1980

TAAAATAGAT AGTGTAGAGA AGAATTTACA AAAAGATATA TCTAGTTTAG ATATCAAAAT 2040

AGATAGTGTA GAAAAGAATT TACAAAAAGA TATATCTAGT TTAAACACCA AAATAGATAG 2100

TGTAGAAAAG AATTTACAAA AAGATATATC TAGTTTAAAC ACCAAAAATAG ATAGCGTAGA 2160

AAAGAGTTTG CAAAAAGATA TATCTAATTT AAACACCAAA ATAGATAGTG TAGAAAAGAG 2220

TTTGAATCAA AAACTTAGCA TGGGTAACAG ACTAGTACAT TTTATGATAA TAACAGCAGC 2280

AATTCTAGGT CCAATTTTAA ATGCCCTATT TATGAGGTAT TTACAATACA TCAAATAATG 2340

ATGTATTGTA TAATTTGATT TTTAAAATGG TACATTATAA TATTGATGAA GAGTATTATT 2400

AATTAACACT TAATTTTTGC TTTTTCATAA AGTAGAACTT ATTTAAATTT TTTAACAAGA 2460

TAACTTAAAT AAGTTCTTTT ATTTTAACAA ATACAAATTG ATTTTAATTC TAAATTGGAC 2520

TATACTCAAT TATTGAAAAG CTTTTTAAAA TTATTTTAAT AAGTGAATTC GGTTAAACCC 2580

TAGCTTTATT AAGTTCTTTA ACAAGAGAAT TTAATAAAGC TTTTATTAAT TATAATAATT 2640

TCTGTAAAAA GTTGGCAAAA AACAATTTAT AATATTATTA TAAATATTAT AGGAGGGATA 2700

TGTTATTATA AATCCGATTT AGTTTGGGCT TAACTAAGTT CTTTTGTTTG AGAATATAGT 2760

TAAGCTCTTT TTTTTATAAA AATTGTTATA AAAAGTTGGT AAAAATAGTT TTTGTTATAT 2820

ATATGTATGT GAATAGCTAA AAAAGTGTAT TGCTATCAAA ACAATCCAAT TAAGTTGGGT 2880

TTAGCTAAGT TCTTAGACAA GAGAATTTAA ATAAGCCCaA CTATTTTTTT TGTAAAGATT 2940

TTTGTAAAAA AgTTGGCAAA AATAGTTTTT GCTATATACn TATATTTATG n 2991

(2) INFORMATION FOR SEQ ID NO: 42:

(i) SEQUENCE CHARACTERISTICS:
(A) LENGTH: 2988 base pairs
(B) TYPE: nucleic acid
(C) STRANDEDNESS: double
(D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 42:

CTATTTGAAA ATTGCGAGCA TAATATGTGT TTTTTTATTT TAGCACAAGT TTTTTAGACT 60

TTCTAAAAAA GTTAAAAAAA AAGAAGATGC TGAGTAATTT GTATAGTTCT TTTAAGATTT 120

TCATTTAAGT AATAAATTAT ATTATCTTGA TATACTTCTA ATATTTACCC ATCAATAAAG 180

CTAGTGTGGC TTAACAAATA AAAACCAATA AATATTTAAA AAATGATTAA TTTAGAATAT 240

AATTTCTATA GCTAAAGCAA AAAATAAATA TAAATTTGGA CTAGTTTTAT TACATAAAAA 300

GATAAAATTA GTCATGCTTG TTGCAATAGA GCCTACTATC TATCCCGCGA GGTATATTTA 360

TTTATATTGC TTTTTAGTTT TTGTAAAGTG ACTTTTAATT ATTAAAATCT AAGGAGAAGA 420

GATTTATGAA CAAAAAATTT TCTATTTCAT TATTATCTAC AATATTAGCC TTCTTGTTAG 480

TATTAGGTTG TGATTTGTCA AGCAATAATG CTGAAAACAA AATGGATGAT ATTTTTAATT 540

TAGAAAAGAA ATACATGGAT AATTCAAATT ATAAATGTTT AAGTAAAAAT GAGGCTATAG 600

TTAAAAATTC TAAAATTAAA TTAGGTGTAA ATAATACTAG AAGTCGTTCT TATTCTTCTA 660

GAGAGACTAA TGTTTCGGAT TCCTATAATA AAACCTATTC ATATTGCAAA AGCAACTGAT 720

TAATTTTATT ACAAAAAAAC AAGAGAATGC TCAACCCATA ATTAGGTGAC AATTAATTGA 780

ATATATGCAG GGATTATTAA AAGTTAGCTT CTGTGACATT ATACACTTGA ATATAATATT 840

ATAAAATAAT AGAATATATG GGTGTTAATA AAGCTTATAA GCATAGAATA TATCATATAA 900

AGAAGAATTT CTATCCCCTT AAAGGAAGTT GATAGTTTAG CTCTTTGTAA TGTCCAACTT 960

GACTTAGACT CTGCGTATAA TGATTTTTTT AGAAAAATTT AAAAGGGAAA TAGAACACAA 1020

GGATTTCCTA AATATAAAAG TAAGAAAAAT AGGGAAACTT ATAGAACTAA TAATCAAAAA 1080

AACTCAATAG GAATAAAAAA TGGTTATATA AAGCTACCTA AAATAGAGTT TATAAAGTTA 1140

TGTCTATAAA TATTATATTT CAATAACAGT TGAGTGCTTA GATACTAAAA ATAATAATGA 1200

AACTAAAGGT GATAAAAAAG AGGCAGTTGG TATTGATATG AGCATGAAAC ATTTTTTAGT 1260

AAGTAGTGAA GGTGAGAAGA TTAATCATCC TAAATATTTA TTAAAAAATG AAAATAAACT 1320

TAAAAAATAC CAAAGAAAAC TATCAAAAAA GCAAAAAGGT TCTATTAATA GAGATAAGTC 1380

TTAAGGTTTA GACTTGCATT TATATCTCCA TCATGCAAAG TTGTTACAAC TACTGAAAGC 1440

CCACCTAGTA TCACTTAATT TTAGAGCCAT ATTTTTAATA TGACAACTAC TACATACATA 1500

TAAATCAAAA TGGTATGGAT CTGCTTTGTA TAAAGTAGAT AGATATTTTC CATCAAGTAA 1560

ACTATGTAAA TTGTCTTGCA AACTCAGACC ATCCTAAATC ATTAATACTT TTTCCAAACA 1620

TTCCTTTTCG CATGCCTTTT TTCATTCTTA GAATTCGAAC TGTAATCAGA CCTTTATATT 1680

CTTTATCAAT TATCATATTT TAGATTTAAA TTGTATATAG CAAAAAGGTC CTATATGGGC 1740

TATTCCGTTT GCAGTAAATA AATAGGGATT GGAATTAAAT CCCTAACCTA ATTGAATAAT 1800

ATCATACTTT CTTCAAGAGG ATTCCCTTCG GCCACTTTTT TTCTTCGTTC TTCCATTACT 1860

TTTTTATATT TTTCAGCTTG TTCTCTTCTT TTTTTATTTA ATTCTGTCGA TTCTCTTTTC 1920

TCTTTTTATA ATTTTTCTTG AACTACTAGT GGCTTAACTG TTTAGATTTG GGATCGTCCT 1980

AGTTATAATT TTGGTACTTC TAATTTTTCT GCTACCACCT TTAATTCTGC CTCTAGATTT 2040

GCTCAAATTT TAGGGATTGA TTCTAATTCA TCTCCTTTTA AAGATTTTTT TCTTCTACTT 2100

TCTTTCCCTT ATGTGCTTTT AAATCAAATC TTTTTGAATT ATCCATAGCT TCTGTTGCTT 2160

TTCATATACT GTTGTATAAT CTAGTGTCAT TTTATTTGGA TCCATTTTAT TTTTAGATGA 2220

TAACTTTTCT AACTTTTTTT ATTATCTTTA TCTTCTTTTT TTAAATCACA CGAAAATAGT 2280

AAAAATAATA GCAAGTAATG GCTAGGCATA TACTTATCTA ATTTAGAGAT TAGCTCCTAT 2340

ATTCAAGCGG CTATTATCCT TATTCTTCTG GCATAGAAGT TGAAAATTTA AATTTTAATA 2400

AATTGTATTT TTATTTTAAT GAGAATAAGC AGAAACATTC CATTCTTAAT TGAATTCATT 2460

AGAAAGTTTC CTTCTATTGC TAATATCATT AATATAATAA AATAATTATC AAAAACATTA 2520

GCAAATCCCC CTTTATCTCT ATGATACTCC TTCACATCTA TATGATTTCT ATCTTTACTT 2580

TCTACATTAG GCTGATTATC TCTACCATAT TTAATATAGC TAAGCGGCTT TTTAACTTTA 2640

CCCATATTTT TCAGTTTGAA TAAAAACCTT TTAACATACT CTTCTATTTG GGATACATCT 2700

CCTTTTCAAT AAAAATTAAA ATGCGCTGAT TTTAATACAT TTACGAAAAA AGTTAATGTA 2760

TCGAGTTTTT CATTACTAAA TCTAAGATTG CTTTTCTAAC TCAGTTTTAA ATTAATACTT 2820

TCATAAGCTT TACAAGCTTT AGTCACTCCT CATATAAAAT CCAAAATTCA ATTGTTTTAT 2880

ATAAGTAAGT ATCAAGTAAA TTTAAATTGG ATAGTAAAAT ATTAAATAGG GGAAAAAACA 2940

AGCTTAATAT TGAGTGATAA ATAAATTTTT CTCTTATTAA ATAGTATA 2988

(2) INFORMATION FOR SEQ ID NO: 43:

(i) SEQUENCE CHARACTERISTICS:
(A) LENGTH: 2970 base pairs
(B) TYPE: nucleic acid
(C) STRANDEDNESS: double
(D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 43:

AGAGTTTTTT CGTTCTTTAA AGTACTTGTT GATTTTCTGG TAACATTCTT TTTTAGGATA 60

CTTTAGCTTA TAGTAAATTT CAGTtCCACA ATTTACACCC ATGTGTTGGT AGTAATTCGT 120

TGTGACTTTT AATACTTTTT CTAATTTATA AAGATAATTT TgCaTTGTTC TCAGTGTAGT 180

GGGAGCCAAA CCATTTCTTT TTAGATTTTT ATTATAGTAA TAGAGTATGT TTTGTTGTGT 240

ATATTTCTTA TCTTTTTTGT TTAGAAAATC TACTGTTGAT GTAAGATATA TTAACTTGTG 300

TTGGTGTTTG TTGTGGCAAG TGGGATTTTT TGTGGTGATT AAAAAATCTT TCATTTTTTA 360

CTCCTTATTT TGTTATTAAC AATTACTATT ATAATGCAAA ATTTTGATTT AAAAGTAAAT 420

ACTTTTCTAA AAAAATATTA AATTTTAATT ATTAATCTTA TTAATTAAAT ACACTTTTTG 480

TAATTTGGTA AAAAGATTTA TTGATTTTAA TCACAAATTA GACTATACTG CAAATAGCGT 540

AGGAAAATAT CTTCATATTT TTACCTACCT TATTTTGTAG TTTTCTAAAA TCATAGTGGG 600

AACTTGGCGA AATTCTTTTT AAAGGGAATT TGGTTAAGTC CCACTTCTTT TGTGTAAAAT 660

TTTTTGTAAA AAAGTTGGCA AAAATAGTTT TTGCTATATA ATTATTTATT ACAAAATAAG 720

GAGGAAAAAG ATGGAAAATC TTTCAAACAA TAATAATCCA CAAGAAAATA TTCAAGGAGA 780

AATTAAATTC AGAAAAGATA TGAGCACCCT AATCAGAAAC TTGCCGCGTA TTGACAAAAG 840

TCTTAAAGGG TATGGGTATA AGTATCAAGA TTTCAATGAC ATAGTAGAAG TAATTTATAG 900

TGTTATTGAT AAGCATAATT TGGATCTTTT TTTTACGCAA GCCCCAATTT CTGTAGAGGG 960

GCAATATGGC ATAGTTGATT ATATTAGGAC TACATTCTAC AGTACAAGCA CTGTGTACAA 1020

ATACTCATTT GATACGCGAA TTCATACAGA TAAATTACAA TGGAACAGTG AAAATGGGTC 1080

TAAAAATATG AATACGATGC CACAATTTGT TGGATCAGCT ATTACTTATT TCAAAAGGTA 1140

CGCTTTAGTA GGGCATCTTT GCATAAGAAG CGAAATGGAT ACTGATGCAG CACCTATTTA 1200

CAATAATTAT GAAAACAGAA ATTCTATGCC TAGCAAACAA TCTAGTGTTA ATCAAAAGCA 1260

AGAACAAAAA AGAGAGCAAA AACAAGAGAT TAATCAAAAT CAAAAAAATA ACACTATTCA 1320

AAACCAGAAA AGAGACATTA AGCAAGAACA AAAAAAAGAT AGGTTTTATT ATTACGGTGT 1380

TTTTAAAGAA GCGTTGTCTA ATATAAAAGA TTGGGTAAAT AGCCCTACAA TAAAAGATAA 1440

TATAAACTCA ATTATTCAAA AAATAAGCTT TATTCAGAAT ATAGACCCCA ATAATGTTGA 1500

TGATATCAAG AAAATTGAAT CTGATTTAAT CTCGTATTTT GAGAAAAATA GTGATTTTAA 1560

AAGTATAAAC TATTGGGCGG AGATTATAAA AAACTATTTC AAGAAAAATA ATAGATTAAA 1620

GGATTTACAA GATTTTGAAA AGTTTGTGTC GTTTAAGAGG ACTGCTTATG GCCCTAGTCC 1680

ATTAATATTC TTTAGTGTCT TAAAAGAATA TGAACGGTTT GATTGCATAT TTGCAGCATA 1740

GCGAATTCTT ATATGGTGAA GCCCCCACAT GGGGGCTGCG ATATTATTGC TGAGCTTGGC 1800

AGGTACTACT TGCACTAGTT GCAAAACTAT CTATACCGCC ACCAAGAGCC CCCTTAACCA 1860

CCTCTTTGAA GGTGCTTTTT TGTTGTTCAG AATTATCCCC AGTACACTTA TCAAGTTCAC 1920

TCTTTATATG ATTAAGTGCA CCTTTTATTT TGTCTTCGTC ATATCCTAAA AATTTATCAA 1980

ATTCTCCAGC ACCAGTTAAA GCGGTTTTTA ACCAGTCAAG ATGTGTTTTT TGGTCTTCAG 2040

ATAGCTTTTC TCTAAGCAGG TCTTCTTTAG ATTTTGGTTT TTCTTGTGTT GCTTCTTTTT 2100

GGGTTAAATC ACGTTTTTGT CTACTTTTTG TTTGGCTAGT ATTAGTATCA TTAGAATTAC 2160

AGCTGTTTAG CATTAGTAAA AATAAACAAA ATAATATGTT GATAATTTTC ATTrTTATTC 2220

CTTTTTTTAT TATTAATATT CACTTAATCA ATTATTAATA CTAAATATTG GATAAACAAT 2280

TATTATTTGA ATTGATATTC TTTAAGTGAG GTAGTAGCTA TTTAGAAATG AAAGCAAATA 2340

TTAGCCCGGC TATCATTGTT ATAGACATTG CTCCCATAAT TCCTAATACC CATTTAAGCA 2400

TTTCTGAAAG AGACATTAAA TTCTTTTCCA CATTGTCTAT TTTAGCAGTA AGTTCATTTT 2460

TAACACTATC TATTTTTAAA TTTAAATTCT TTTCTACAGT ATCTATCTTA GTATCTAAAC 2520

TATCTATTTT TAGATTTAAA TTCTTTTCCA CATTGTCAAT CTTAGTATTA AGTTCGCTTT 2580

TAACAGCATC AATCTTAACA TTTAAATTCT TTTCTACAGT ATCTATTTTA GAAATAAGAT 2640

TATCAAATTT TATATCAAAT TGTTTTTCTA AATTTTCTAA ATCTCTATAT GTTAGTTCAT 2700

TGTGATAATA TCTTTTAGAT AAATCTTGTG CTATTAGTTG TTCCATGCCC AGTCTAATAA 2760

ATTCTTTATA TATTTGTTCT TGAGTTACAC CTGCAATATT TGTTGACACT GTTTCCATAA 2820

AATTTTCCCT TATGGTCATA TTATACACTA TTTTAGATTA ATTGGCTTTA GAGATTTTTA 2880

TATGTAAAAT AGAATTTCTT GCAAGAAAAA CCTTTTTGTA ATTTACATTT TTAATTGGGA 2940

ATATTTATTA TAGACTTTTT CCGCTATTGG 2970

(2) INFORMATION FOR SEQ ID NO: 44:

(i) SEQUENCE CHARACTERISTICS:
(A) LENGTH: 2942 base pairs
(B) TYPE: nucleic acid
(C) STRANDEDNESS: double
(D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 44:

AGAATAGCTT GAAGTAAGGA TAAATGAAGT AGAAAATTTT GCCTGCTTAG TGCAAATTTG 60

TCTATTCCTA ATAGCAATAA CACTTCTGAA TGGAGTTTTG TTACTAAAAG TTCTTCTTCT 120

CCCGCAACTT GTGCTTGTAA GCTTTCTTGT TCACTCATTT TCACTTACCT TATACTTTTA 180

ACTTTGCTTT ATGTTAACTT GCAAAATAGT TTTTCTAGTA GCAAGTAGAC CGCCTAAAAC 240

AAAATCAATG TATGAATGAG CTACATCGGT TGAGTCTTTA TCAACTTGTT CATTCGGTGT 300

AGGTAGCATA TACTTACTAG GTTTAAATTT AATAAGTTCT GAATTGAGTG GATAAATGAG 360

TATTTTATGT TTTAGCAAGT TTGAAGTTTC AATGTAAACA TCTTCTCTAT TATTAATAGC 420

CTTAATAGTT TGAATTAAAA CATCTTCCCA TTTTTCACAA CTACTTGCTG CACCCTGTGC 480

TGCTGCGTAT GGTTTTACTA GTTTAAGTGA CGTTGCTGGG TCAACTATTA CCATCATCGG 540

TGTGGAAAAT TCGTCGCCCA GTTCCAACTT TGAAAGTCCA GCCTCAATTT TTTCAAATAT 600

TTTATCCATT TTATCTTTAT CACCACTAGC AACCTCTTCT TTTATCTGCT CGGGCATATT 660

AAGCAGCCCA TACATATTTG GAAGCAGGCG TTTTTGATTT TTTCCATCTT TTTGAATCGA 720

AACAGTTCCT GTTAGTACAA AGTGATTAAT AAGTTTAATA ATTTCACTAC TTGCAAGCTT 780

ATACGCTTGA GCAAAAGGAA GTAAATTATT ATTAATGTCC CCAATATATG AATCTGAAGT 840

ATAAAATTTC TCAGAAGTCT GCTTTAAATG TCTGAATTTA TACTGTAATT TCAAATAATT 900

AAGTCTTACC ACTTCAGAAC TAAATCCAAT AGTTGAGATA GTATTAACCT CATTGGCAAT 960

TGTTGTAGGA TTAGCATTTA AAAACGCGTC CCATTTTACG GTTTTTTGAT ATCCCATTTG 1020

TAGATCAACA TCTTCAATTT GATCGGGCGA AAACCATTTA TACATAATAG GATCTTTAAC 1080

TTCTCCTATG ATATTTGCCA CAGCTTTTGC ATAATAATTT TCATCAAATA ATTCCATATT 1140

AAATCCTCCT AAATATTATT AATTTCTACT CATAGCTTTA TTCCCAAATA CAGCTACTTT 1200

TACTAAATAA ACCTCATTGC TAATTTGTTT CGCATCAGTC AATGCTGTTG CATTAATAGT 1260

TGCTTTATTT GGTGCTCCAG TCACCTTTTC AAGAGCACCG TCTTTATTAA AAACAAGTTT 1320

GTCTTTTACT TTAACCGTAG AATCTTTTGC CACTAAATAT CCCTCAAAAT TATTGGTAAT 1380

TGGCACAATA GTAGCTGTTT TACTAAACTC ATCTATGTCA ATGCATATTC CATATAAGTC 1440

TTCACCTCCA CCAGCCTCAA CATGTGGCTC ATAGTGACTT TGATTATCTT GAGCCTCTTG 1500

AATAACTCTT TTTACTCCAC GCTTGTACGG ATACCCCAAA AATGGATGAT TTTCCAATTT 1560

ATCAAACTTA CTAGTTCTAG TGCCTCCAGA AGCAAAAAAT TtCaCATTTT TGTCTCTAAA 1620

TTCATTAGAA TTGCTAAGCA AACTAGCGTC ATGTTGGGGA TTTTtCATAA ACTTTTCCAG 1680

TTTACTTCTT TtCTCTkGAT ACTCTTTTAC TAATTGCGTT GTATCTCCCA TTTATTTACC 1740

TCCTTTTATT CGCCCAAAGT TTAACCACCA TCAGGTATTA CTATCTTCTC AAGGCCTCTA 1800

TTTCCAAAAA TTGCAACTTT TATCAAATTA ATAGAATACT CTTGCCTAGG ATATCTATTT 1860

TGATCTtGAT TTCCAtCTTC GGGTGCAAAA TTGATtGTAA ATGAATCAGA TAGAGCATAT 1920

ATGTTAATTA CGGTTGGTGG CCCACCTCCA GCCTTGATAA TAACACCGTT ATTATTTATG 1980

TCTAGGATTT CTCCTATTTT TATACTTGGA TTCCTTGTGA CAAGGTACCC TTCAAAATTG 2040

TTAGTAATTG GCAATACATA CGCGGTACAA CTAAACTCAC ATACATCTAC ACATATCCCA 2100

TACATATCAG TATCAGCTCC AACTTCyACA TATATAGAGT TCTCTTTTGG AACAAGTTTA 2160

ACCCCACGCT TGTATGGAAA ACTATTTGCT GGGTCGTAAA GGTATTCCTC TATTTTGTCT 2220

GTATAACTTG AACATGCAAA TGAATATGCA TCAACTCGCT CATTCTTAGA TTTAAAACAA 2280

CTACTCAAAC CGCCAAAAAC CTTATTTTCA ATTGAACTCA TAGATTTTAC ATATTTCTTG 2340

AATTTCAAAA GGATATCATC AAGCTCGTTA ATTGCCTCCA AATAGGGATC TTCCCCTTGT 2400

GCATCCTCAG CTTGTCTTGC TTGCCGTTTA GCTCTAGGAG CAGCGGAAAC TTGTGCCCCT 2460

AAATCTACCT GTGGGTCCTC AACAGCCTCA AGATTTTCTA CTTGCATGTT GCCTTTTAAA 2520

GCCATAATTT ATTACCCTTT TATTGCTCTA TTCCCAAATA CACTAGCAAG CACTATAAAT 2580

AAATCTTCAG TTAATTTGTG TGCCTTTGAA AGTGCTATTG CATTAACAGA TTTTTGAGCC 2640

CCAGTGACCT TTTCAAGTTC CCCATGTTGG TTAAAATTTA ATTTATCTCC TGGATTTACA 2700

CCATTTTGTC CTTCTTTCTT AAGCGTTAAA TACCCAGTAA AGTTATTTGT AATTGGTATA 2760

ACAGTTGCCA TACCGCTAAA CTCATCTATA TCGGAACACA CTCCATATAA ATCGTCTCCA 2820

CCACCAGCCT CAACTTCTAG TTCGGTTGTT CCATCTCCAA AACTAAGCTT AACACCCCGT 2880

TTATACGGAT ACCCTTTAGC AGGGTAATTC TCTATTTTGT CTTTACTGCT AGTGCAAACC 2940

CC 2942

(2) INFORMATION FOR SEQ ID NO: 45:

(i) SEQUENCE CHARACTERISTICS:
(A) LENGTH: 2892 base pairs
(B) TYPE: nucleic acid

(C) STRANDEDNESS: double
(D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 45:

GGGTCTATAA TGTCATTTAT TATTTCCTCA GTGCTTTTTC CAGTTTTTAT TCCGTATTCT 60

TGCGCTGCTT TATAAGAGCT ATACATTTGA CTTTTTTGTT CTTGCGTTGC TTGTTCGGTT 120

TCATAAAGTT TATTTATTGA CTTTTTTAAA TCTTCACTAA GATTATCATA AAAATTTGAA 180

ATTTCATTAG TATGCATATT AATTATAGAT AGTATATAAA TAAACAATAT TTTGAGCAAT 240

AGTTTTTGGC ATTTTTTAAA TGAAAGTTTT GATAGAAAAC ATTTCTATAT TCATAACAAT 300

GAAATCTACA AAAAAATAAC AGCCAGTGAA CTTTTCTACT AGCTGTTACT TTGTATACGC 360

AAATTTAGTT ACACCTAAAA GCATCCAATA TAATTACTGA CTGTCACTGA TGTATCCTTT 420

AATTTCTTCA AATTTAGAAC TATCTTTAAG ATATTTTTTA ACTTCTTCTA ATTTTGATTT 480

TAATTTTTCT AAATCTTCTT TAATTTCGCT AACATTTACA CTTTCTTTTA ACTTAGGCTC 540

TTCGTAACCA GTATATGGTT TATTGCCTTC ATTTAATTTA GTTCTTAGCG CGTCCCTAGC 600

ATCACTCAAT TCTTTCAATA ATTTTCCTAA TCCTTCGTCT TCTGATTCCT CTTCTAACCC 660

CTCCCAAGTT TCGCGTATAG AATTATTGCC ATTGGTAAAA TCATCATACA CAGGCCCAGT 720

AATTCTATCT ATAACGCCTT GTGGGCCTAC AGTCGTTTGC CATTTTATAA CATCAATATT 780

TTCATTGAtC TCATCTATTT TAGCTATAAG TGTTTTAATT TGGTTATCAA CTTGCCTTTT 840

TTCTTCCTCT TCTTGTTGTC GTTTTTGTTG TTCTGCCTCT TCTCTTTCTC TTTTTTCTTT 900

TTCTGCTTTA GCTTTTCTTT GTTGCTCTTC TTGATGTCTT TTTTGTTCTT TTTGTTGTTC 960

TTCTTCTTTA GCTTTAGCTA ATTCTTGTTT GTTTTGTTCG TCTATTATTT CTTTTTCTTT 1020

AATGTTTTCT TCTGTATTCT CTTGTTTTTC TTTATTTTCT TCAACTTTAG CTTCAGCTTC 1080

TTCTACTTTT TTTTCTTGTT GACCATCACT TTGTTCTATT GCTTTTAATA CTAATGCATT 1140

ATTGTGAATA TTTTCCGGCA ATACTGGTGG CGGATTTATT CCACTGCCAT TAGGATCATC 1200

GCCCTGCATT AATTCTTCTT TTTCTTCTTC TTGTAATTTT TTTGCTACTT CATCTACTTT 1260

TGTACCACTT GAAGCAATTT TATCCTTTAC TGGATCTAAA ATCTTATTTA CAAATCCTTT 1320

AATTTTCCCT TCTGAATTTT GTTTTATATC TTTACCAGTT GCAAAATTCT TGCAAGAAAT 1380

TATCAGCGCA AAAACAGCAC AAATAATCAA TGTTTTTTTA TTCATAATTA TTCTCTCCTA 1440

TATTTCTAAA TTCTATTTTA AATTTTTTTT AAGCACTTCT AGTGGTATTG CATATTCAGC 1500

TGTTTTATCT CCCCCCACAC CGTTATTAAT TTTATCTCCT AAAAACGCAA TATGTTCAGC 1560

ACCATTAATA CTTTCTATTT TTGTTATGAT CTCTTTGTTT TGGATACCAT TTGCACTACT 1620

TTGTTCGTAC TTATATCCAT AATACAAACT AGTTTTAAAT GATCCGCCTT TAGTCATTGC 1680

TTTTATAAAG TTATTAACTT CTGATTCTTT TAATGAAAAG AATGTAGCCG AATGTCCCCC 1740

AGCGTTTAAA CCCGTATCAA TACCATTTTC TTCTTTTCTT ACAACTAAAT CTCCTAGGTC 1800

TGTCCAGTTA CTATTATTAT CTTTATTTTT AATTTTTACA GTAAATTTAG AGAATTCTAT 1860

TTTTTTAAGT TTTAACTCAC CATTACTTTG CTCATCATAT AAAGTATGGT TTATCTCACC 1920

ACTACTTTGC TCATCATATG AAGTATGAAT TTTGCAAGCA CCTATAAGTA TAAAAACAGC 1980

ACAAACAATA AACATTTTCA TTTTCTTATT CATAAATTTC TCCATAAGTC CTAATCATAC 2040

CACAACAGCT AATAATTGCA ATATTTCAAA GATTTAAATA TATAATTTTG TTACATTCAG 2100

CTATTACATA TTAACAAAAC TCAAATGTAA TTTTAACCAA CTCCCAAAAA TCTCTCCATT 2160

GCAAATGCCC GGCTCATTAC AAAAGACTAC AAAACACATA CAAATTAAAT TTCAAAGTCT 2220

TTGCTATATA TCACTTAAAG TATCATGTCT TTCTTAAGTC CACCCCTTAA AAATTGCCTC 2280

TTCTGTTTAT CACAGCCACT CCACAACCCA AATTTCGCAT GCAATGAGAA CACCATAAAT 2340

TTGACTAAAA TTTTAGGGTT TTGATAAAAT ATAAATTACA TTTTTATTAA ATTTTTATTA 2400

CTTTTACTTA ATTTAAAAGT AACACTTCAA AGGAGAGGAT TTTATGGATA CTAATAATTA 2460

TTTTAATTTA AATAATTTCG ATACAGATTT GATGCACAAA TTCTTAAAAG ACTATCAAAA 2520

TGTATTAAAT GAAAACAAAA TTCTTAAAAA TTCACTAAAA ATTTCTTCTA AGCCTACTAA 2580

AAAAGCTTCA AAACCAACTC CAAAGTTTTA TTTGAATCAA AAAATTATCA AAATAATTGA 2640

AAAATGTGTT AAAACATTAA AATAAATTGA CCCAATTTCT GGTTGGTTTT TAAATCTACT 2700

GGCAATAAGT GGCTGCAGAG GTGCCGAGCT GCAAAAAGTA AAAATGCAAG ATATTACTCC 2760

CTTTTTAAGC AAAACTGGAG AAACTTTTTA CAATATAAAA GTAAATGTAG CTAAAAAAAG 2820

AAATGTCACT TGCATTAGAG AAATTGTCAT AAAATCTGTA GAATTTGATG CTATTCAAAA 2880

AGCTCACGAA AA 2892

(2) INFORMATION FOR SEQ ID NO: 46:

(i) SEQUENCE CHARACTERISTICS:
(A) LENGTH: 2849 base pairs
(B) TYPE: nucleic acid
(C) STRANDEDNESS: double
(D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 46:

GCTAGAAAAG CATATGCATT AGCAAGGGCA TATTATATGG AATTTAGATT TAAAGCCGGA 60

AGTGTTTTTT GCTATTTTAA AGGGCTATAT CGTTTATTGG ATAAAAAAAG AACAAATAAT 120

CATTACAACA AAGTTTTATT TAGTATGTTT ACGGATTTAG AACAACAAGT ATATAAATTT 180

TATGGGAAAA AATACCCGGA ACAAGGACCG TTAATAAAAT GGATAATAAA AAACCTAAAA 240

TAATAACAAT AGCGTCAtTA AGGGCGGTGT TGGCAAAgCA CAAGTGCAAT TATATTTACA 300

ACTCTATTGT CTCAaGATtG GAAAGTGCTT TtAATTGATA TGGATACACA GGCATCAGTA 360

ACTAGTTATT TTTACAAAAA AATAATAGAA GATAATTTTA ATTTATTGGA AAAAAATATA 420

TATGAAGTTT TGAAGGGGAA TGTATTAATT GATAATTCAG TTATAAATAT TAGTAATAAT 480

TTAGACTTGA TACCTAGTTA TATAAGTTTA CACAAATTTA ATAAAGAGGC TATAACATTT 540

AAGGAAATTA AACTTCAAAA ACAGCTATTA AATTTACAAT CCAATTATGA TTACATAATA 600

ATTGATACAA ATCCCAGCCT AGATTATACG CTAACCAATG CTTTAGTATG TAGTGATTAT 660

ATAATAGTTC CAATAACAGC AGAGAAATGG GCTGTTGAAA GTTTAGAACT TTTAAAGTTC 720

TCAATTAGTG ATTTAGCCAT TGATATTCCA ATTTTTTTAA TAATAACTAG ATTTAAAAAA 780

AATAATACCC ATAAGGCACT ATTTAGTTCG CTTAAAGACA ATAAGAATTT TTTGGGGTTA 840

ATTTATGAAA GAGAAGATTT AAATAAAAAG ATAGCAAAAA ATGATCTATT TAATTTAAAT 900

AGAGATTATA TGCTAGAGTA CAAAAATATA TTAAGTAAAT TTATAACAAT AATCATGTCC 960

AGGTAACTGG ACATGATTGC CATTTTAAAT GAAAGGAGTC CATATATGGA GATAATATTG 1020

AACAAAAGAA ACCTAGAAGT GCTAAATGAA GCCGAAGAAC ATTACAAGAA GTTAAaGCAA 1080

AGATTAAAAT CTAGTTTTCA ACAAGAAATT TATTATAAGA TGGAAGTTAT TAAGATATTA 1140

AAAGAAATAA AAGATAACGA ATATTATAAA TTAGATGGAT ACAGAACATT TGAAGATTTT 1200

ATCAAAGATT ATCATTTAGC AAGGAGTCAA GCATATGACT ATTTGAAAAT AGCAAATGCA 1260

ATTAAAGATG GCATTTTAGA AGAAGCTTAT GTAATAGAAA ATGGTGTTAC AAAAACTCTT 1320

GAGTTCTTAA GAAAATCGCC AAATGTTTTG AAAAAATCTA AACAAAATCC AATAAAACCC 1380

TTAAGATTTC AACTTAAGAG TCAAGAAAGT TACGACTTTT ATAAAAGTAA TGCTAAATTT 1440

ACTGGATATC TTTTAGACAA ATTATTTAAT AATGAAAAGG AAATGATTAA AAAAATTATG 1500

AAGGAATATA AACAACTGAa AGGATaGTAA GAAGTTTTAT GACTAATTTA GCGTACAGAA 1560

CATATAACAT AGAAAGCATA AAAAATGAAT TTTTAAACAT AGGGTTTAGT GAGGAGGCAA 1620

TAGATTTTGT TTTTCTTCAT AATGATAATT TCAATTTTGA ATTTTTAArA GAGAAAATAA 1680

TCGATTTAGA AAAGAATTTG AGAAAAGATA TATCTAATTT AGATATAAAA ATAGATACTG 1740

TAGAAAAAAG TTTAAATCTA AAAATArATA CTATAGAAAA AAGTTTAAAT CTAAAAATAG 1800

ATTTTGTAGA AAAGAGTTTA AATGCCAAAA TAGATAGTTT AGATACCAAG ATAAATAATG 1860

TAGAAAAAAC TTTACAAAAA GATATATCCA GTTTAGATAC TAAAATAGAT AGTGTAAAAA 1920

ACGAACTTAA TTCTAAAATA GATAGTATAG AAAAAACCTT GCAAAAGGAT ATATCTAGTC 1980

TAGATAATAA AATAGATGTT TTAAAAAATG AACTTAATGC AAGCAATAGA ACAATACAAG 2040

TAATTCTAAT AATGGGAATA ACACTtGCTC CAATTATCTA TTCTATATTT AATAAGTATT 2100

TCTTTAATTG AGAATGATTA AAATTTTTTA AAATATTAAG GGAGTATATA GCGTATTTTT 2160

TAAATAGAAT ACTATAATCT TGATTTAAAT TCTTTAAAgA AACATTTTAT TTTTACTTTC 2220

TTTTAAATTT AGAACTTATT TGAATTTTTT AACAAGAAAA TCTAAATAAG TTCTTTTATT 2280

TTAACAAATA CAAATTGATT TTAATTCTAA ATTAAACTAT ATTCAATTGT TGAAAAGCGT 2340

TTATTTATTA TAATAATTTC TGTAAAAAGC CTGACAAAAA TAGTTTTTGT TATATATATG 2400

TATGTGTATA GCTAAATAAG TATATTGCTA TCAAAAAAAT CCAATTAAGT TGGGTTTAGC 2460

TAAGTTCTCT AACAAGAGAA TTTAAATAAG CCCAACTAAT TTTTTGTAAA ATTTTTTGTA 2520

AAAAAGTTGG CAAAAATAGT TTTTGCTATA TACTTATATT TATTACTATA AAAGGAGTAA 2580

AAAGATGGAA AATCTTTCAA ACAATAATAA TCCACAAGAA AATATTCAAG GAGAGCTCAA 2640

AATGATAAGT ATTAATCAAC AAAGTTTTAC TGGTTGTGAA ATATTTGAGG AAAAATCTTC 2700

TCCCATTrAA GAAAAAAGTA AATTAAGTAA GATAGGCAAG AAATTGCCAG GaATAAGTAG 2760

TCAAGAATGT TTTAGATTTA ATCGCAATAT TGATTTTAGT GTGCAAAGAA ACAAGTTAGA 2820

TAAATACGGT GCTAGTGAAG TAGGCAATA 2849

(2) INFORMATION FOR SEQ ID NO: 47:

(i) SEQUENCE CHARACTERISTICS:
(A) LENGTH: 2682 base pairs
(B) TYPE: nucleic acid
(C) STRANDEDNESS: double
(D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 47:

AGTTGCGTAT CTTGACATAA AAAGTGAATT TGATACTGAT GCAGCACCTA TTTACAATAA 60

TCACGAAAAC GAAAATTCTA TGTCTAACAA GCAAGTTAGT GTTAATCAAA AACAAGAACA 120

AAAAAAGGAC ATAAATCAAG AAAAAAATCA ACTGAACACC TTTAATAAAA ACTTAAAATC 180

TGGCAAGGCT TATTGCTATG AAATTTTTAG AGACGCACTG TTTAATATAA AAAATTGGGT 240

AAATGAAGGT GAAGAAAAAA ATAATATAAA TGCTCTTATT CGGGCATTAT GTACTGATAA 300

TGATGATGCT TTAGAAGATC TTTTTGAAAA GAATGCTGAG CTTAAGAGTA TAGAATATTG 360

GGTAAATTTT TTAAAAAAAT ATTTCAATAA AACTAATAGA TTTGATGATC TAAATAAGCT 420

TAAAGTATTT ATGTCTGATA ATCGGGATGT TTATAAAACA AAAGTATTAA AATTCTTTTG 480

TATGTTGAAA AAAGAAAGAC AATTTAATTA TATATTTGCA GCATAGCAAT ATTAAAGCCC 540

CCTATTTGGG GGCTGCTATA GGTATAATAA TTAAGGTTTT TATTTTTGAT TGAGAAATGT 600

TTTTAGTTTT GCCAATTAGC TGTAAGACCA GCATAATGAT TATCTTCTTT TAAAAgTTCA 660

yCTTTAAgGT ACtGATAAAT TTCTTCATTA GAATTTCTAT TAGaCATATC ATTAGCGACG 720

CCTCTAAAAT aTTGCyCTAy TAAGTTGTCC ACCTCCCCAT ATTTCATTAT TTTTATTACA 780

ATCTTGTGGA GTGTTTTTTT TACAATCAAT AGCTCCTTTA ATATAAGTAT CAAAAGTTTC 840

ATTTTGTGCT TTTGATTTTA AGAAGTTGTA AACCTTGGTA AAAGCACCAG CTAATTCTTT 900

TTGTTTTTGA ATATCTTCAG AAAGCCAATC AAAGAAGTCA TTACATTTAC TTTTGTTTCC 960

ATTCATGCAC CCTTGTATTT CATTGTTTAA TTTTTCAATT GTGTATTTAA ACACATTGAT 1020

TAATGAAGTA AACATTTTTT CTTCGTCGGA TGTTAAAGTG ATTTTTTCTT GTTGTGGCAG 1080

TTCTTCTTGG CTTAAATCAC GTTTTTTCCT GCTTTTTGTT TGTTGGGCAT TGTTTTTTAA 1140

AGTGTCATTA TCATTGGAAT TACAGCTATT TAGTAGTAGC AAAGATATAC AAAATAATAT 1200

GTTGATGATT TTCATTGTTA CTCCTTTTTT TATTATTAAT ATTCACTTAA TCAATTATTA 1260

ATACTAAATA TGGGATAAAC AATTATTATT TGAATTGATA TGTTTTAAGT GAGGTAGTAG 1320

CTATTTAGAA ATGAAAGCAA ATATTAGCCC GGCTATCATT GTGATAGACA TTGCTCCCAT 1380

GATTCCTAAT ACCCATTTAA GCATTTCTGT AAGAGACATT AAATTCTTTT CTACATTATC 1440

TATTTTAGCA TCTAAATTAG ATATGTCTTT TTGTAAATTC TTTTCTACAT TGTCTATTTT 1500

AGTATTAAGT TCGCTTTTAA CAGTATCAAT CTTAACATTT AAATTCTTCT CTACAkTATC 1560

AATCTTAGTA TCTAAATTAG ATATATCyTT TTGTAAATTC TTTTCTACAk TATCwATCTT 1620

AGTATCTAAA yTAgaTmTAT cTTTTwGtaA ATTCTTTTCy ACATTrTCTA TCTTGGTATT 1680

AAGTTCACTT TTAACAGCAT CAATCTTAAC ATTTAAATTC TTTTCTACAG TATCTATTTT 1740

AGAAACAAGA TTATCAAATT TTATATCAAA TTGTTTTTCT AAATTTTCTA AATCTCTATA 1800

TGTTAGTTCA TTGTGATAAT ATCTTTTAGA TAAATCTTGT GCTATTAATT GTTCCATGCC 1860

CAGTCTAATA AATTCTTTAT ATATTTGTTC TTGAGTTACA CTTGCAATAT TTGTTGACAC 1920

TGTTTCCATA AAATTTTCCC TTATGGTCAT ATTATATACT ATTTTAGATT AATTGGCTTT 1980

AGAGATTTTT ATATGTAAAG TAGAATTTCT TGCAAGAAAA ACCTTTTTGT AATTTACATT 2040

TTTAACTTCA GATATCAGTT TTAAATTTTT TACTGTAGAT TTTTTACAAA AACAGTATTG 2100

CAAAAACTCT TAGATTACTT TTTCTTTTCT TTGTATACTA CAATAACTCC AAAACCCACT 2160

AAATGGTTTA GTGATTTAAC CTCAAGAATA TTTTCGGCAC CTATTTGGTT AATAAAATTT 2220

TCTAACCCTA TCCCTATAAT TTCGAATAGA GTTTTGTTTT TATCTTCTTT TTTTATAGGA 2280

AAGTTAATGk TATGCTTATG ATCATCACCG CCTTGATCTA AAGCaTTAAA GTTTTAACTT 2340

TaTAATTTCA TCyTTTTaAT TcATATGAAA TTAAATTACC AATACTGATA ATAAACATAA 2400

aTAACATTAA TAAATTAATT TTTTGcACAT tGTGTTCCTT AATAAATAGA ATATTAACAA 2460

TATTATATCT TTATTAAGAT TTGCCCTAAA ATATAAAATT TTATTAAAAT ATAGCAGTAA 2520

TAAACGACTT TAAGAATATA AATGGGAATT TCTTGCAAGA AAAACCTTTT TGTAATTTAC 2580

ATTTTTAATT GAGAATATTT ATTATAGACT TTTTCCGCTA TTGGTTTTGT TTTTTTAATG 2640

TACTCTAAAT ATATGTTGAG GGTACTCTGA GCTCAGATTT TT 2682

(2) INFORMATION FOR SEQ ID NO: 48:

(i) SEQUENCE CHARACTERISTICS:
(A) LENGTH: 2532 base pairs
(B) TYPE: nucleic acid
(C) STRANDEDNESS: double
(D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 48:

CGGCATnACC AGTAAAGTTC AACACTAGAA TATTGCCTTA AATGCACGTA TAAATCCTAA 60

TTTTAAATTA GCACACTCTA TATCTAATTC ACTTATAACT TTCCtAGCGT TAACTTCTGA 120

TTTAAAAGTT TGTGATAAAA GGTGTTCTAA AGTATCTTCA CTAATTGTTA CTCTAGAGTC 180

TTGGTTAACA ACACTTTCTC CACTTTCCCA TTTTTTCCTC ATCCTCCACA CATTTACCCT 240

AGAAACCCCC AATTTATCCG CTATTTCCCT ATCATCTAAC AATCCTTCTC TAAAATATGC 300

AACATAATCA TCAAAAGACC TTTTGGCTCT TTTCAAGAAA ATTCTCCTAA AATAACAAAA 360

TTAACAAATT GTTACTCTAA ATAGTAAAGC AATTTGTTAA TTCAATTAAC ATAAATTATT 420

AATTTCTTTA TACCTATTAA CAACTGCCCT ATATTTACAA TATTTATTAA TATAAAAACC 480

AAACATTTCA AATATCCAAA AAGGAGCATT TATATGAATC AAAAACAAAT ATTTTTATTA 540

TTTTTATTAT TTTTAAAAGT AACAATAAGT TTTTCTTATG ATCAATCTCA ATACAAGGGA 600

TATATGGAAA AATATTATCA TAAGAAAGGC AAAACAGATA CGCACATATC CTTTTTTCAA 660

ACTCTTAGTG CTGATGAAGG GGGTTTTTCT ACTATCTTTA TAGGAGAAGA TGAACAATTA 720

AGTCGTCTTA GTTTTACCAC TTTAAAGGAT ATTAAAGACG GCAAAGAAAC CTCTTATATG 780

GGTTTTAACT TAGAATACCA TTACAAAGCA AAGTTTAACA ATCCATACCC TATGTTAAAT 840

GATATACGTG CAAACATTAG TAAAGTGAAA GTTAAATTTT TTTTTGATAA TGGCCCCGAA 900

AAAATAATAA GAGAATTAAA TCAAAAATTT GTAAATAATA GAGTTATGTG GGAAATTTGG 960

AATAATTCAT ATAATAAGCT TTCAGAATAT ATTAGGATTA ATCTAAGAAC CTCTGATCCG 1020

GGCATAGAAA ATTTACTGCC AAAATTATTA AAACATAAAA CCGTAACAAT AACAATTGAA 1080

ATTCCTGAAA GTGAAGATCC TGAGAAATTA ACTAGTTCTA TAACTTTTGA TCTTGATAGC 1140

TTTCAAAAAC TATACAAGAA ATATAGTACG TATTTTAAAT AATATAACAA GACTCGCTGT 1200

GAGTCTTGTT ATATTATAAT TTATGATTGT AAAAAAAATT TTTCTATTCT TTTTTATTAG 1260

AATCTTTAGA TTCTTTCTCT AACTTTTTAA GCTCTTCAAG CTCTTCTTCA ATCTTTTTAA 1320

GCGAGCTTAC TATTATTTCT TTAGCTATAT CGCTAGTACT ATTACCACTA GAAATATTTT 1380

TAAAGCCCAA ACCCCGAGCA TGCCGCAAGG nTTCTATTCC AACTTTCCCT TGATTTTTAG 1440

CCCTTTCTCC ATCAGTTATT CCAATTGCAG ATTCAACTTG ACCCTTAAGT TCTTCAAATT 1500

TTTTTTGAGA CTCTTCTAAT TCTTTTTTTC TTTTCTCTAT TTTTTCATTT AAAGCTTTCT 1560

CAAGGATTTC TAATTCTTTT TCAAACTTTT CTTTATCTTT TAGTTTTTCC TTTAATTCTT 1620

CTATTTCCTT TTCATAATCA GAATATGTTT TAAGAGAAAC ACTTTTAGGA TCCGACTTCT 1680

CTATCTTATC CTTTAATTCT TTTATTTTTT GTTCAATTTC TTCTTTTAAT TTTTGATCTT 1740

CAGCAACACC TTGAACCACT TGATCTTCGC CCTGCTCCTG AGGCTCATCA GCTTGCATAG 1800

ATTCTTCATC CTTTGGTTGA ACTTTTGCTT CTGGTTTTTT AAAATCTCCA AAAAACTCTT 1860

CTTTTTTTGT ATCTAAAAAT CCCTTAACTT TTCCTTTTAA ATCTTGTTCT AAACTTTTTT 1920

TTACATCTTC ACCACTTGCG TAATTCTTGC AAGAAACTAT CAGCGCAAAA ATAGCACAAA 1980

TAATAAACAT TTTCTTATTC ATAAGTTGCT CCATAAGTCT TAAATCTAAC GCAACACCGA 2040

ATAATTACAA TTTTTCAAAG ATTTAAATAT ATAATTTTGT TACATTCAGC TATTACATAT 2100

TAACAAAACG CAAATATAAT TTTAACCAAC TCCCCAAAAT CTCTCCATTG CAAATGCACC 2160

ACTCATTACA AAAGACTACA AAATCCATAC AACTTAAATT TCAAAGTCTT TGCTATATAT 2220

TAGATAAAGT ATACTGTCTT TCTTATCCGA CACCCTCAAA AAATGCCTAT TCTGTTTATC 2280

ACAGCCACTC CACAACCCAA ATTTCGCATG CAATGAGAAC ACCCAAAATT TGACTAAAAT 2340

TTTAGGTTTT TGATAAAATA TAAATTACAT TTTTATTAAA TTTTTATTAC TTTTACTTAA 2400

TTTAAAAGTA ACACTTCTAA GGAGAGGATT TTATAGATAT GAATAATTAT TTTAATTTAA 2460

ATAATTTCAA TATGGATTTT ATGCTCAAAC TATTTCAAGA TTATCAAAAT GTGGTAAATG 2520

AAAATAAAAT TC 2532

(2) INFORMATION FOR SEQ ID NO: 49:

(i) SEQUENCE CHARACTERISTICS:
(A) LENGTH: 2528 base pairs
(B) TYPE: nucleic acid
(C) STRANDEDNESS: double
(D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 49:

nAGGATCCCC AGCTCCCttA ATAGCCCCCa TCAGCCTTCC CTTTCTCACC ACCATCGTTC 60

ACAGCAAACT TTCCATCCTT AGCCATCCCC CTCAAAGCAA TAGCAGCAGC AATCTGATCA 120

TCCTTCTTCA TCTCATCCTT AAACTCCGCA CCATTCTCCT CATTACCCTT CCCAATAGCA 180

GCAGCAATCG GATTTGTAGC ATCCCCAGGC TTCTTTCCCT CCTGATCAGC CGCATCAGCA 240

GCCTTAACAA TCGCACTTAA TATCTGCTCC CCACTAACAG CACTAACAGC ACCAGCCGCC 300

TTGCTAGCAG CCTCACTGTC CCCAGCATTA GCACCAGCAC CAGCCTTCCC AAACAACTTC 360

CCTGCCCCTT TATTACTCTC CCCTGTAGCA GCAGCAACTT TCAGCTTTTC ACTCCCCCCA 420

GCAGCTTCAA CAATCTCCTT TATCCCCTTA GCAATCCCCG TCACACTCGC CTTATCAGCA 480

ACCTTTGCAG CACCAGCATT AGCCACAACT TCTCCAATTG CATCAGTACC ACTTGAAGCC 540

CCCTCAGCTG TCTTTACAGC TTTTACCAGC TTATCCAACA ACTCAGCAGC TCCCTTAATA 600

GCCCCCCTCAG CCTTCCCTTT CTCACCACCA cTCtTCACAG CAAACTTTCC ATCCTTAGCC 660

ATCCCCCTCA AAGCAATAGC AGCAGCAATC TGATCATCCT TCTTCATCcn TCATsmTTAA 720

ACTCCGCACC ATTCTCCkCA TyACCCTTCC CAATAGCAGC AGCAATCGGA TTTkTAGCAT 780

CCCCAGGCTT CTTTCCCTCC TGATCAGCCG CAtCAGCAGC CTTAACAATC GCACTTAATA 840

TCTGCTCCCC ACTAACAGCA CTAACAGCAC CAGCCGCCTT GCTAGCAGCC TCACTGTCCC 900

CAGCATTAGC ACCAGCACCA GCCTTCCCAA ACAACTTCCC TGCCCCTTTA TTACTCTCCC 960

CTGTAGCAGC AGCAACTTTC AGCTTTTCAC TCCCCCCAGC AGCTTCAACA ATCTCCTTTA 1020

TCCCCTTAGC AATCCCCGTC ACACTCGCCT TATCAGCAAC CTTTGCAGCA CCAGCATTAG 1080

CCACAACTTC TCCAATTGCA TCAGTACCAC TTGAAGCCCC CTCAGCTGTC TTTACAGCTT 1140

TTACCAGCTT ATCCAACAAC TCAGCAGCTC CCTTAATAGC CCCCTCAGCC TTCCCTTTCT 1200

CACCACCACC ACTCTTCACA GCAAACTTTC CATCCTTAGC CATCCCCCTC AAAGCAATAG 1260

CAGCAGCAAT CTGATCATCC TTCTTCATCC CATCATGATT AAACTCCGCA CCATTCTCCG 1320

CATCACCCTT CCCAATAGCA GCAGCAATCG GATTTTTAGC ATCCCCAGGC TTCTTTCCCT 1380

CCTGATCACC AGCAGCCGCA CCAGCAGCCT TAACAATCGC ACTTAATATC TGCTCCCCAC 1440

TAACAGCACT AACAGCACCA GCCGCCTTGC TAGCAGCCTC ACTGTCCCCA GCATGAGCAG 1500

CATCAACTTT CCCAAACAAC TTCCCTGCCT TTTCATTGCC CTCTTTAGCA GCAGCAACTT 1560

TCAGCTTTTT ACTCCCCCCA GCAGCTTCAA CAATCTCCTT TATCCCCTTA GCAATCCCCT 1620

TCACACTCGC CTTATCAGCA ACCTTCGCAG CATTATCATC AGCCACAACT TCTCCAATTG 1680

CAGCAGTACC ACTTGAAGCC CCCTCAGCTG TCTTTACAGC TTTTACCAGC TTATCCAACA 1740

ACTCGCCAGC TCCCTTAATA GCCCCCTCAG CCTTCCCTTT CTCATCATTC TTCACAGCAA 1800

ACTTTCCATC CTTAGCCATC CCCCTCAAAG CAATAGCAGC AGCAATCTGA TCATCCTTCT 1860

TCATCTCATC CTTAAACTCC GCACCATCCT CATTACCCTT CCCAATAGCA GCAGCAATCG 1920

GATTTTTAGC CTCCCCAGGC TTCTCTCCAT CCTGCGCAGC CTCACCAGCA GCCTTAACAA 1980

TCGCACTTAA TATCTGCTCC CCACTAACAG CACTAACAGC ACCAGCCGCC TTGCTAGCAG 2040

CCTCACTGTC CCCAGCATTA CCAGCACCAG CCTTCCCAAA CAACTTCCCT GCCTTTTCAT 2100

TATTCTCCCC TTCAGCAGCA GCAACTTTCA GCTTTTCACT CCCCCCAGCA GCTTCAACAA 2160

TCTCCTTTAT TCCCTTAGCA ATCCCCGTCA CACTCGCCTT ATCAGCAACT TGGCTTTTAC 2220

AATTAATAAA AACAAAGAAA GTTGTTAATA AAATTGCACT TGAAATTTTT TTCATATTTT 2280

TTTGTTTAAT GATTGTTTTG AACATTTAAA AAATGTTTTT GTTAAGAGGC TTTTATTCTT 2340

TGTTTAAGTT AAAGTTAAAT AATAATAACT AGTTTTTTTA ATTGGATTTG TGAATTTTGC 2400

CTACTTCCGT ATCACATATA ATAAAGTTTG ATGACATATA TAGTATTTTT ATTGCTTGAG 2460

AGTTTTAACA AGTGTAGGTG AGAGAATTTT TCATTTCTTT TATTATAAAG AATAAAAGAA 2520

ATGAAAAA 2528

(2) INFORMATION FOR SEQ ID NO: 50:

(i) SEQUENCE CHARACTERISTICS:
(A) LENGTH: 2496 base pairs
(B) TYPE: nucleic acid
(C) STRANDEDNESS: double
(D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 50:

TGCAGGTCGA CTCTAGAGGA TCCCTGCCTC TTyaAGTATG CTAGCCAAAA TATraCGAGT 60

AAwTTCTTTG TCTGATACTT TAAATTCTTT GTCATATATA TTTTTTkcAA TTTTAAATAC 120

TATAGArTCA TCAGGcTcTC ATAAAGyATC TCTCTAAGAG TTTTTTGAAT TATmTCTTTT 180

TCtTTrgAtA TTTGyTCTTT TTCAACTGmT ATTAyATTrC TTGTyTTTAG GTATCTTTCT 240

TTTCTTATAA GATAGCTAAA TAAAATAAAC CATTCTTGCT CTTTATATTC TTCTTTtTCT 300

TTAATTTTAG AAAAATCAAA TTCAATATAT CTTTTTTCTC CATAAAGTAC TTTCGATTTA 360

TCATATAATC TCCATACCTT TCCATTTGAA AGTATCCCAT AATGTTTTTG ATATTGATTT 420

AGATATCTGT ATAGCTGATC TTCTGCTTCT TTTACCTTAT CTTTAGCATC AAAACTAAAT 480

GATGGACGCT TAACTTCTGC TATAAGCAAG ATATCTTCAG TTGGAATAGA TTCATTATTT 540

TTTTTAGCTT CTTCTAATTT TTTATTAAAA GCTACTTTAT CTTTATCATT TTCAAAAAGT 600

AGTATATCTA CTCTGGATTT TACTCCTTCT ATTTGCCCAC CTTTTTGTTG TTCTACTGAA 660

TAAGCTAATT CTTCAAATAT AGACTTTAGC AAAGACTCTA TATTTGCTTC TGTTGAATTG 720

TCATCTATAG CTTGAATTTT ATTTTTTAAA AAAATAAAAA AGTTTTTTGA TTTAACAATA 780

TTTTCTTTTT TTATAAAGCC TTTTGACAGT TCTTTATAAA GAGATACATT TGGATCATTT 840

GTTTTTATAA TGAATCGGCT TTCATTGTTC ATATTTACAA CCATTATGTT ATTTATAAAT 900

CCTTTTTAGC CCTTCTTGAT ACTCGAAATG TACTCTAAGA TTAGTTTTTT TAAAATTAAA 960

AAAACTAATC TTAGAGTAAG TCGGCCAAAA CTTGTTTTAA TATTTATTTG ACTATCAATA 1020

CTCTATCTTT AGAATAAGCT TGTTAAAAAT TATTTCTCAC TTTTTCTCAA TTAATTAAAG 1080

TTATTAATTT ATTTTTTATA AGGCATCCTT AATTAAAGAG CATTTAAAAA ACACTTTTTT 1140

AAAACCGAAT TTTTCTTAAA CATTCCCCAA TTTGTGAAGC ATAAACAAAA AAATGTTTTT 1200

ATCCTTTTCA TTTTCAAAAT TACAATTATA GAGTCTTTTG TTAATTTCTT CTTTAAAAAC 1260

ATCTTGCTCA GAATCATGCA AGCAACAAAG ATGCAAAAAA TTTTTAAAAG AACTTATCAG 1320

GTCAAAAACA ACACAAATAA TAACCCAGTT TTTATTCATA ATTATCCTCT CTCAAAATTA 1380

AAAAATAAAT CAAAGTCTTT GATTTACAAT TTTTTATTAA GCAGTGAAGG GGGAAAAACA 1440

AATTGTCCTA AATATTTAAC AAAAAATGGA AATAGACTTA ATAAACTTGG TTTTTTCTCA 1500

GGAAGGATTT CTAATTACAA CATCAAATTC TTCCTGAATA TCTGGAAACT CAATTCCATA 1560

GTAAAAAAAT TTATAGTAGC CGCTTAATTT TCTAATTATA TCTATTTTTT CTTTATCTTT 1620

ACTAGAAGTT TTATTGCCTA AAATATTTTC AATTAAACTA ATTAAAGCTG TAATATCACT 1680

AAATTTTATA TTTAAAGATT TGTCAAAAGA TAATGAATAA AGTTTAATTA AAGAAAATAT 1740

TATTCCTAAA TTATCTGTAT CTTCACTTTT CTCATATTCT TTATATATAT TTTTAGATCT 1800

TTCTATATCT TCTTTAGCAG TATCGTTAAT ACCTCTAATT TTTTGATAGC TATATTCTAA 1860

AATAATTGTA ATTTCTTTAA TTCTCTTTTT AAATAAAGAA AATCCACCTT CAAATTTCTT 1920

TTTTTTAATA TCAAAAAATT AATCTTTGGC ATATCCTAGT AAAGGATTTC CTACTTTTAT 1980

ATGATGTTCA ACAAAGCTTA ATGGTGTTCC AAAAATAAAA TATTTTAATA ATCTAACTTA 2040

TTTTTTTGAG ATAGTTTTTT ATAAAACTTG CCCAAGCAAT CTATACTCTT AAGTTCGGGT 2100

TTTCTCAATC ACCTATAACT TTATTTGCTA ATTAATTTTT TATAAAAGCT ATCCTTAAAT 2160

TTTTCAAAAT ATACTATATG AACTACTGCT TAAAAAGCAA AGACTATAAA ATAAGTAGTT 2220

CATCAGAAAG TTTTTGATGG TATTACTACT ATTAATAGAA TTTAAAAAAT CGAGCTTACC 2280

AATGTATTTA AATAAACTAG AAAGCAAACC ACAATACATG ATTAAAATGA TATGGGCAAT 2340

AAACTTAAAA AGTTTAACGA TAATTTTGAA ATTCTTATTA GAATCTAATT AAAAATGTAA 2400

ATTTATATAA ATTTTATAAA TAAAGAGTCA AAGAAAACGC TTTATACTAG AAAGTCTTTA 2460

TAAAGATAAT AAAATATATT TTTTAGAACT TTCTTC 2496

(2) INFORMATION FOR SEQ ID NO: 51:

(i) SEQUENCE CHARACTERISTICS:
(A) LENGTH: 2487 base pairs
(B) TYPE: nucleic acid
(C) STRANDEDNESS: double
(D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 51:

ATTTATTTAC ACGAAACCCT GACTTTTTGA GAAGATTTTT GAATTTATTA GAAATATGGA 60

TAATGCTAAT TTGGTTGTCT TTAAACTTAT GTTTGGATTT TTGAAAAAAA TAAGTACGTC 120

TTGTATCAAG ATTTTTCTCA TTAAAATGAT TTTTGTGTGC TGTTTGGATA GCCTCGAACT 180

CTTCTGAGTT GATAACAATT TCTCTAATAC AAGTTACATT TCTTTTTTTT GCCACATTTA 240

CTTTTATGTT GTATAAAGTT TTTCCATTTT TGCTTAAAAA AGTTGAAATA TCTTGCATTT 300

TTACTTTTTG CAGTTCGGTG CCCCTGCAGC CACTTATTGC GAGTAAATGT AAAAACCAAC 360

CAGATATTGG ATCAGCTTGT TTAAGAGTTT TGATGCATTT TTCAATTAGT TTGCCAATTT 420

TTGGGGTCAA ATAAAATTTA GGAGTTGGCT TTGAAGCTTT TTTAGTAGGC TTAGAAGAAA 480

TTTTTAGTGA ATTTTTAAGA ATTTTGTTTT CATTTATTAG TTTTTGATGA TCTTGTAATA 540

ATTTAAGCAT AAAATCTATG TTGAAATTAT TTAAATTAAG ATAATTATTC ATGTCCATAA 600

AATCCCCTCC TTATAAGTGT TACTTTTAAA TTAAGTAAAA GTAATAAAAA TTGATTAAAA 660

ATGTAATTTA TATTTTACCA AAAACAAAAA AATTTAGTCA AATTGTGTGG CTTCTCATTG 720

CATGCAAAAT TTGGATTGTA GGATAGCTGT GATAAACAGA AGAGGCAATT TTTAAGGGGT 780

GCACTTAAGA AAGATACTAT ACTTTAAGTG ATATATAGCA AAGACTTTGA AATTTAAGTT 840

GTATGTGTTT TGTAGTCTTT TATAATGAGC AGGCCATTTG CAATGGAGAG ATTTTAGGGA 900

GTTGATTAAA ATTATATTTG CGTTTTGTTA ATATGTAATA GCTGAATGTA ACAAAATTAT 960

ATATTTAAAT CTTTGAAAAA TTGTAATTGT TTGGGGTTGT GGTAAACTTA AGGCTTATGG 1020

AGTGGATTAT GAATAAAAAA ATGAAAATAT TTATTATTTG TGCTGTATTT GTGCTGATAA 1080

GTTCTTGCAA GATTGATGCA ACTGGTAAAG ATGCAACTGG TAAAGATGCA ACTGGTAAAG 1140

ATGCAACTGG TAAAGATGCA ACTGGTAAAA ATGCAGAACA AAATATAAAA GGGAAAGTTC 1200

AAGGATTTTT AGAAAAGATT TTAGATCCAG TAAAGGATAA AATTGCTTCA AATGGTCCAA 1260

TAGCAGATGA ATTGGCAAAA AAATTACAAG AAGAAGAAAA GGTAAATAAC GGGGAAGAAG 1320

AAAATGATAA AGCTGTCTTT TTAGGAGAAG AATCAAAAGA GGATGAAGAA GAAAATGAGC 1380

AAGCTGTTAA TTTAGAAGAA AAAAATGCGG AAGAGGATAA GAAAGTTGTT AATTTAGAAG 1440

AGAAAGAATT AGAAGTTAAA AAAGAGACTG AAGAAGATGA AGATAAAGAA GAAATAGAGA 1500

AACAAAAACA AGAAGTGGAA AAAGCACAAG AAAGAAAACA ACGACAAGAA GAAAAGAAAC 1560

GAAAAAAACA AGAACAGCAA GAAGAAAAGA AACGAAAACG ACAAGAACAA AGAAAAGAAA 1620

GGAGAGCTAA AAACAAAATT AAAAAACTTG CGGATAAAAT AGATGAGATA AGTTGGAATA 1680

TTGATGGTAT AGAAAGTCAA ACAAGTGTAA AACCGAAAGC AGTTATAGAT AAAATTACGG 1740

GGCCTGTATA TGATTATTTT ACCGATGACA ACAAAAAAGC TATATATAAA ACATGGGGAG 1800

ATTTAGAAGA TGAAGAAGGC GAAGGATTGG GAAAATTATT GAAAGAATTG AGTGATACTA 1860

GAGATGAGTT AAGAACCAAA TTAAATAAAG ATAATAAAAA ATATTATGCC CATGAAAATG 1920

AGCCTCCTCT AAAAGAAAAT GTAGATGTCA GCGAAATTAA AGAAGATTTA GAAAAAGTAA 1980

AATCAGGATT AGAAAAGGTT AAAGAATATC TTAAAGACAA TTCTAAATTT GAAGAAATTA 2040

AAGGATACAT CAGTTACAGT CAGTAATTAT ATTGGATGCT TTTAGATGTA ACTAAATTTT 2100

ACGTACACAA AATAACAGCT AGTAGAAAAG TTCACTGGCT GTTATTTTTT TGTAGATTTC 2160

ATTGTTATGA ATATAGAAAT GTTTTCTATC AAAACTTTCA TTTAAAAAGT GCAAAAACTA 2220

TTGCTAAAAA TGTTGTTTAT TTATATACTC TCTAGAGCTA TGACGTATAC AAATGAGATT 2280

TCAGATTTTG ATGATAATTT ATATAAGAAA ACAAAAAAAG AAATAGATAA ACTTATAAAC 2340

AAGCTCTATT TAACTAGCCn AATAACTCTA AAGCAAAAAA GACAAATnTA CAGTGCTGTT 2400

GAAAGAATGC AAAAATACGT AATAAAAACC GGAAAAAGTG TTCTTTTAGA ATCGGAAAAA 2460

GAATTTGTTA AAGACACTTT GAAAAGA 2487

(2) INFORMATION FOR SEQ ID NO: 52:

(i) SEQUENCE CHARACTERISTICS:
(A) LENGTH: 2462 base pairs
(B) TYPE: nucleic acid

(C) STRANDEDNESS: double
(D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 52:

AGGAGATAAG TTTATTGGTA TTTTTTATGG CTATAGAAAC CCAATCAAAA cCTTTAATAA 60

AGTATaAAAT AAATGGAACT AGAAAAaGCAT ATGCATTAGC AAGAGCaTAT TATATGGAAT 120

TTAGATTTAA AGCCgGAAGT GTTTTTTGCT ATTTTAAGGG GCTATATCGT TTATTGGATA 180

AAAAAAGAAC AAtAATCaTT ACAACAAAGT TTTATTTAGT ATGTTTGCaG ATTTAGaACA 240

ACAAGTATAT AAATTTTATG GGAAAAmATA CCCGGAGCaA GGACCGTTAA TAAAATGGAT 300

AATAAAAAAC CTAAAATAAT AACAATAGCG TCAATCAAGG GCGGTGTTGG TAAAAGCACA 360

AGTGCCTTAT TTTATGGCAA TATTTTAGCT AAAGAAAGAC ATAAAGTATT GATAATTGAT 420

AGTGATCCAC AGGCCAGTAT TACTAGTTAC TTTTTGTTTA AATTAAAAGA ACAAAATGTG 480

AATGTCGAAA ATTACAATCT TTATGAAGTT TTTAAACAAA GAAAATATAT AGAAAATTGC 540

ATTTTTACAG TATCTAATTG TTTAGATATA ATTCCCAGTT CCTTAGAATT ATCTGTTTTT 600

AATTCAGAAA GCATACCATT ACAAGACAAC CTTTTAGAAA AAAGACTTTT GACTATTAAA 660

TCTAAATATG ATTATGTGAT AATCGATACA AATCCCAGCT TAGGACATCT TTtAAACAAT 720

GCTTTAGTAA TTACCAATTA TTTAATAATA CCAATTAATT CCGATTTATG GGCAGTTGAA 780

AGTATAGATC TAATATTAGA TGCAATAAAT AAAGTTTATA GAAATGATAT TACACCTTAT 840

TTTTTAGTGA CGGGGGCACT AGAGAGACAA AACATAGATA AGGAAATAAT ATTTAATTTG 900

GAGAATAGAT ATAAAGAAAA TCTAATAGGA GTTATTCCTA AAAGAGATGA TATCAAAAAA 960

GTGCTGTTTT ATAGAAAAGA ATTTTCTTCA AAAACAGACT ATTATCAAGA ATATAAAAAA 1020

TCTTTAGATA AAATGTTAAA AATAAAATAA CAAATAAAAT ATATCCAGTA ATGGACAAAT 1080

AAGGAGTTTG CATGAGCATT AAAAATAAAA TGATAATAAC CAAAAGAATA GATATAAAGG 1140

AAAATATGTC TAAAATGGAG TCATTAGAAG AAATTCATAA AGAAGAATAT TTGAGATTAA 1200

AAGACAAATT AAAAACTCTA ACAACGGATG ATATTTATAA TAAAATAGAA ACAGCAAAAA 1260

TATTAAATGC GATTAATCAA AAAAAACTGT ATATTTTAGA CGGATATAAA AATTTTTATA 1320

GCTTTTTAGC TGATTTTAAA ATCGCTAAAT CTCAAGCATA TAAATATATA AAAATAGTAT 1380

CGGGCGTAGA AAAAGGTATT ATTGACTATA ATTTTATTGC TAATAATGGC ATTGAAAAAA 1440

CAATTAAACA ATTGGAAAGT AACAATGTTA TTAAAAAATC TAGGCAAAAT CCAATAAAGC 1500

CTTTAAGGTT TCAACTTAAA AAGCAAGAAA GTTATGATTT TTATAAAAAA AATGGGAAGT 1560

TTACTGGGTT TTTATTGGAA GAACTTCTTG AAAGTCAAAC AGATTTGATT AATAAGCTTT 1620

TAAAAAAATA TAAACAATTA AAAGGATATT AAGAGGATTT TATGAGAAAT TTGGTGCACA 1680

GAACATATGA TATAGAAAGC ATAAAAAATG AATTTTTAAA CATAGGATTT AGTGAAGAGG 1740

CAATAGATTT TGTTTTTCTT CATAATGATA ATTACAACTA TGAGGTTTTA AAAGAGAAAA 1800

TAATTGATGT AGAAAAGAAT TTGCAAAAAG ACATATCTAG TTTAGATACT AAGATAGATA 1860

ATGTAGAAAA GAATTTAAAT GTTAAGATAG ATAATGTAGA GAAGAATTTA AATATTAAAA 1920

TAGATAGTGT TAAAAATGAA CTTAATTCTA AAATAGATAG TTTAGATACT AAGATAGATA 1980

ATGTAGAAAA AACTTTGCAA AAAGATATAT CTAGCTTGAA TACTAAAATA GATAGTGTAG 2040

AAAAAACCTT ACAAAAGGAT ATATTTAGCC TAGATAATAA AATAAATGTT TTAAAAAACG 2100

AACTTACTGC AAGTAATAGA ACAATACAAG TAATTTTAAT AATGGGAATA ACGCTkGCTC 2160

CAATTATTTA TTCTATATTC AATAAGCaTT TTTTAAATTA AGAaTGATTA rAaTTTTATA 2220

AaGTAATAAG TTAGTATATA GCTTTtAAAGT AGAACTTATT TGAATTTTTT AACAAGAGAA 2280

TTTAAATAGG TTCTTTTATT TTAACAAATA CAAAATAATT TTAATTCTAA ATTGAACTGA 2340

ATTTAATTGT TTAGTGAGTT TATCTAAAAT AAATTGAGCT AAGCCAGCGG CTTTCTTAAG 2400

CTCTTTAACA TGAGAATTTA ATAAAGCTTT TATTTATTAT AATAATTTCT GTAAAAAGCn 2460

TG 2462

(2) INFORMATION FOR SEQ ID NO: 53:

(i) SEQUENCE CHARACTERISTICS:
(A) LENGTH: 2447 base pairs
(B) TYPE: nucleic acid
(C) STRANDEDNESS: double
(D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 53:

AACCCCnAAA AAGGGCTCCC AATATTAAtw GAGAGATTTA TATTTTTCrA ATGTTGTGCT 60

AGCTTTTATy TCATTATTAT TGAATATAGG AGTAACTAAT GAGAAATAAA AACATATTTA 120

AATTATTTTT TGCAkCAATG TTATTTGTAA TGGCTTGTAA AGCATATGTA GAAGAAAAGA 180

AAGAAATAGA TTCATTAATG GAGGATGTTT TAGCTCTTGT AAATGATTCT TCTGGAGGCA 240

AATTTAAAGA TTATAAAGAC AAAATAAATG AATTAAAAGA AAATTTAAAA GATATAGGCA 300

ATGCrGAGCT TAAAGAAAAA CTATTAAATT TGCAAAATTC CTTTCAGGAT AAATTAGCGG 360

CCAAATTAGC AGCGTTAAAA GCAGCTAAAA ATACCATTGA AAACATTACT GACAAGGATC 420

AGGATATTTC AAAAAGAAAA ATATGGTCAG AAGCAAAATT AGTTGGAGTA ACTGTACCAC 480

TTCTTGGAAG CAATACTTCT GGTAATGGGG ATAAAATGTC TAAAAACGCT GTAGAACAGA 540

TAGACAAAGT AATAAAGTTC CTCGAAGAnG GCACTAATTA ATTAGCAAAT ATTCCTGTTG 600

AAAATGTTAA AATCTAGATA TTAAATCTGC GTCAATCTAA TATCTAGATT TTTCTTTTGT 660

TGCAAAAGCC GATTTGATCA TAAATTAGAA TTTCTTGCAA GAAAAACCTT TTTGTAATTT 720

ACATTTTTAA CTTCGAATAT TGATGATATA CTTTTTCCGC TATTGGTTTT GCTTTTTTAA 780

TGTACTCTAA ATATATCTTG ATGTTATGTT TTACCGCAGt ATAGAGTGTT CGTCTTTTAG 840

TGTTGATAAG TCTGGATAAG GATATTCTGG ATAATTGGGG TCATTAACTT TCACTTTTGT 900

TTTAGCTAAA AATGTTACAA GATACATAAC ATATTCTGAA AGTTGTGTTT CATATTTAGC 960

TAAAGATTTT AGCGTTGGAA TAATTGGCGG TTTTGGTTCT TCTGGTAGGT TAGCAATAGT 1020

GGTGCAACAT AACAACAAAA CAATTAGTAA ACAATGCAAT CTTTTAAGCA TTTTCACTCC 1080

TTTTAAGCAT TTTGATGTAT TCTTTCATAA TTTTGTTGCG TTTTGCTTTA AGTGAAGTGA 1140

TAATTTTTTT ATTTTTGTCA TAAGAGATAG CCTCTATTAT CTCAATATTG TATTTTAAAA 1200

TGTCTTTTAT TTCTTCAAAT ATTTTAGTTG AyTCAGCTGT TTCCATAGAT TTTAGTGTGC 1260

TTATATATGT TTTATAGAAA AAATCTATTA CTTTACTAAA AGTATTAATG TAATTATGAT 1320

CTATGTTTGT ATCAGTTTTA GCTATGTTAG TTAAGCTAGA TAGTAAATTA AGTCCCAATT 1380

CAATAGTGTT CTTTTGCATT ATTTGCTCTC TCTTTGTAGA TAGGTTTTCC GTCTTTATTG 1440

AATTTTAGAT CATTGGATAT TTTTAGATTT TTATCATTAG AATTAACCAA ATCAATAACC 1500

GTATTGATTT TTGCATTTAA AGGAGCGAGT GCCGCATTTA TTGCTGGGGT TAATGCACTC 1560

TCAAGTCTTT CCATATTTGC TGTATAGATT AATTTATAAT GAGAATACAG CTCATAAACC 1620

AAAAAGAATC CTTTATGTGC AATTTCATCA AATTCATCTT CAAATTTAGA AAATATATCA 1680

ATAAGGGTTG ATAAAGACGT AAGTCCAAGC TCAACATTAT CTTTGGATAA TTTCATAAGT 1740

TAATCTCTTT TTTTAATGTG ATTTTTGCCA TTACCATTAC CATTCTTAAA AATCTTGCCT 1800

ATTACAATAG TCAATATGTC TTTTAGTAAA GGCTTGAGAA GAATTAATAC TCCTAAAACC 1860

AACACTGTTA CAAAAATCAT TACGGCTATA AGTTTAATTT CGTTTAAATT GATAAGAAGT 1920

TCTGTTAATT TAATAGTATC CATTTTTTAA TCCTTTATTT TAATTTTTTA TTTGTATATA 1980

CATTATATAT CAAAATCGTA ATTTTTGCTA AAAAAGCTTA CAGTTTTAAA AGATCTGGGA 2040

CTGAATCTCC ATACATGTAG GCTCTTTTTT GGATATACCA TCCGTTATAA ATGGGAGTTC 2100

TGGTGAATAG TCCGCGAGGG CCCGTAATTT CATAAACCCA ATTGTATGTA ATTTCATCAG 2160

AATACATAAG TTTTAATGAT TTTTTCTGCA TCCTATTATC ATTGATTTTT ACTTGAATCT 2220

CAATATCAAG ATAAATGGGA TCATTATCGT AATTTTTGTA AAAAATAAGA GTTACATCAT 2280

CATTGTCTTC CATATTAATA GATATCGCTT TATTCTCATA GTCAGAAGGA TAAAGTGTAG 2340

TACTTCTAGT TATGCTGCCA TAAGTAAAAT TTGATGGTAC TCCAAGTATA TGTTTAGGAA 2400

CnGGTGTTTT TTGAATAGTA TCTGATGAAG GCATTATCAA AAGATCA 2447

(2) INFORMATION FOR SEQ ID NO: 54:

(i) SEQUENCE CHARACTERISTICS:
(A) LENGTH: 2401 base pairs
(B) TYPE: nucleic acid
(C) STRANDEDNESS: double
(D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 54:

TATTATTATA TTTAGGGAGA GAATTTATAA AATAAAGCAT AATTACTAGA ATACTTGTTT 60

TTTCTATGAA TGATTGTAAT GCGGATCTTA ATATTAATAA TTATAAAACC AAAACTAAAG 120

ATGGGTTTTA TGCTTTAATT TAATTAATTT TAAGGAAAAA CTAATTACCA TATTAATCCC 180

AAATATATAA TTATTGACAA AAGTTGTATC AAGGGATATT GCCTAATATA GCAGAAGTAT 240

ATTCTGTTAT TAAATATCTC CTAAGGAGGA TTTTATTTTT AAAATAATAG AAAATAGTGC 300

ACTTATTTTA ATAGATATAC AAAATGATTT TTTAGAATCA GGCACTTTGC CAGTATCTAA 360

CAGTAATGAA ATAATTTCTT TGATTAACCA ACTTCAAAAT TATTTCAAAA ACATTATTGC 420

CACCAAGGAT TGGCATTGTA AAAATCATGT AAGCTTTTCT AACAATAAAA ATGGGGGTAT 480

TTGGCCTGAG CACTGCGTCA AAAATACTTG GGGATCAGAA TTTCCTAATG ATCTAAATAC 540

GAAAAGAATA AAAAAAGTTT TTTTTAAAGG AACCGATCAA TATTACGATA GTTACAGTGG 600

ATTTTATGAT GATTGCATTA AAAAAAAACA AACGGGCCTT CAGCTTTATC TGAAAAACAA 660

TTCAATCAAT ACATTATTTA TAACGGGACT AGCATTGGAT TTTTGTGTAA AAGAAACAAT 720

ACTTGATGCA ATTAACTTGG GATTTCGAGT TTATCTAATA ACAGATGCTA CAAGAAGCAT 780

AACATCTACT CCTGAATTAA TAATTCAGGA ACTTAAAAAG CTTAATGTAT TAACTTGCTT 840

CTCCAAGGAC ATCTTCGACA GCCAAAGTAA GCTTAATATA TAAAAAATCA TTCAATAGTA 900

TTTAATTAGA AAACTACTAT TTATAATTAA AACTATCATG GAATGATAGT TTTTTAGACT 960

ATATAAGAAA AGTTTATTCA CCAAAGAATG GCCTTTATAT TAAATTAAAG CCGCCTTTTC 1020

CTTGGTTTTT ACTTCTTAGT AAGAATAATT TTAAGATTCA TAGTTACATT TATATCTCTA 1080

TCATATAAAG CTCTGCAATT AACACAAGTC AACTTAATAT TACTTATCCT TTGTGTAATA 1140

CCACTTCGAA TGCCCTATTT AAATATAATA AAAGAATGAT AAAGAACTCA AATACTTCCC 1200

CCATGCATCA TATAAAAATA TCCATCTCAA AATGTTAAAA ACAACGCCAC CGACATAATT 1260

CTCTAATACC CAATGTACAG TCTATAGATA CAATAAAATC TTTTAAGAAT TTAATTCCAA 1320

TAACACCTGT TTAAACCGGC TCACAATATA AAAAATTTTT ATTTTTTAGT GATAAACTTT 1380

ACTTATCTGA AAAATTGCTT TATTATGTAA GAGTGTATAA AAAACCATCC GAAGTTGAGG 1440

AGGCAGAAGT GAAGGTTAAT AAATCCCTAC AAATACAAAG TAAATATCAA CACAAACTAA 1500

TTGCTTTAAT TGCGACACTT GAGTATATTA ATAAAAACAA AAAAAAATAC AACCAATCAG 1560

ACATCCTTTA TTGTTTTAAC AGTAACTTAA GGCGCAACGG GCAAAAAGAA GTTTCAATCA 1620

AAACGCTTAG AAACTACTTC TATAAACTAG AAAAGCTAAA TATTACTATT AACTACTATA 1680

GACATCTAGG TATTAATATG GGCACTGAAA TCTACTATGC TCTTAGGCAT TCTAAAAAAG 1740

ACTGCTATAA TCTACTAAAC CAACACTTTA GGAATAAAAA AACAGAAAGA TTTCAAAGAC 1800

GTGTTAATGT ATATATTAAA ATAAATTACG ATAAAAAGGA CAATGTAAAA AATGGGGAGT 1860

GTCTTAATAA TAAATATAAA AAAGAAGAAA GAGAAACCGA AAGAAAAAAA AGGATTAATA 1920

AGCTTAAACT AAAAAAATAT GCAAAAAAAT GTAATTTCGA TAATGAAATT TCCTCTTTTA 1980

TTATTAATCT TAACTTAAAA AAAGAAACAA CAATCAAGCT TTTTAAATTT ATAATCAAAG 2040

AAAAATATTA TTTCAAAAAA GAAAACAAAT GTAATTTACA AAAAACACTG CAAAACAAAA 2100

AAAGAGATTT AATTTCAATA TTAAGAAAAA CCCAAAAAAT TTTAATAAAA GAAGGTTGCG 2160

ACAAAAAAAA GATAAAAACC CAAATACAAA ACACATATCA AAAATATAAA AACAAACCCC 2220

ATTTCATATT AGAAAGCAAT AAATATAAGG ATTTCGATCA AATTATAAAA AAGATAAAGG 2280

ACGATACTAA TAAAACCGAA CCCCAAAAAC ATAAAGACAA TATAGAAACC AATATATATA 2340

ACATACTTTT AGATCAATTA CATAGAAAAA CCAACACAAC AAATTTAAGG TCGGGGATCC 2400

G 2401

(2) INFORMATION FOR SEQ ID NO: 55:

(i) SEQUENCE CHARACTERISTICS:
(A) LENGTH: 2324 base pairs
(B) TYPE: nucleic acid
(C) STRANDEDNESS: double
(D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 55:

CAATATGACA TCCTGAGGAC CTTTTATGGA GACTTGTAAA GCTTTTTATT TCTAATGTTT 60

TAGCTTTTAT AAAAACACCA ACCATGACAC ATATTGAATT TTTAAATCTA TAAACATTAT 120

TTCCTTAAAT TTCTAAAAGT TTTTTAGGCT CTGTATTTAA AAAAATCACT TCACCAAGAA 180

TAACCTTTTC ATCATTTAAT AATAATGTTT TCTTGCTAAG AAAATTTATA AATCTATTTA 240

AAATGCTTAA TTAAGCTTAT TTGTTTTCAA ATAATTCTCA TATCCTTTTA TTAAAAACAA 300

AATGTATTCT TCTCCCTTTT TATTTTTTAG CACCTCAAAA TCATTAAGCA AAACCTCAAA 360

ATCTTCTTTG GTTAGCGAAT AAAGACTAGC TACAATAAAG TTATTTTCAT TTTCTTTTTC 420

TTTGAAAAAT TCATCTTTAG TGTCTAATTT TAGAATTTTA TTAACTTTTT CTTTGCTAAA 480

TTTAAAATGC TCTAAGTAAA GCAAATTAGA GAAATTTAAA GGATCATTTT TAGCTATTAA 540

CAAGGAAGTG TTTTTTACTA AAGTTAAGTA TATCGGATTA GCTAAAATTT CTTCTTCTTC 600

GGGTTGAGGC ATAGGGCATT GATATAAGCA TGATTTTACA ATATCAGTGC TTAAAGCAAA 660

TCTTCTTATT AAATAGTCAA AAACAAATGA ATTAAAAATA GATATAATAA ATAATTTTTT 720

ATAAATAGAT ATTGGTGTTT TCTCATAATT TATATATATT GTAGAAACAC AATAACAATT 780

TCTAGGAGAT AAAGTACTAA TCATGGTTCT TATATCTGTA TTTCTTGCAA TCCTTCTATA 840

TAATATTTTT TCTATTTGAT ACTGATTGTC TTTAGTTGAT ACTTTTTGAA AGTCATCTTT 900

ATCTATCCAT AGTAATTTAG AGCTTTCTTT TGCATCTTTG TCTTCAAAAA ATCTTGAATT 960

AAACTGATGA ATATTAGCTC CAGAATAAAG AAATATAAAA TTTTCATTAT TATATTCTTT 1020

ACATAGTGTT TTATACTTTG TTAAATTTAG CCCTACTCCA AAATTAATAT ATTCTTCACT 1080

AAGAGTACTA AATTTGCTAA ACATTTTGTT AATTAAGATA AGCTCTTTAC TATCTTTAAA 1140

TTCAATAATT GATTCTTGAA TAGGAGACAG TTTTTTAATT TGCTCTATAT CTAATTTAAT 1200

TCCTTTATAA GGATCATCTT TATTATTTTC TAAGTTACTG GTTATTTCTT TTAAAATATT 1260

ATCATTACTC TGAATCATAA ATTTTGCTTT AAAATTCGAT GTAGGAGTTT TAGTATTGCT 1320

TATTTGAAAT ATTGCAAATT TAAAAAGTGT TGCCACATCT TTAAATCTTT TTTGATTTTG 1380

AAATTGATAA ATATAGTTAA GCTTATAGTT AGTAAATATA TATTTTCTTA GTATCCTAGC 1440

ACTAGATTCA CTCCAAAGAG CTGAAGGAAC TAAATAGGTT AAATTTCCGT TTTCTTTTAT 1500

TAATTTCAAA TTAAATGCTA CAAAATATCT AAAAAGATTT GGATCACCAC CACTAGCAAA 1560

ATTTTTAAAA TCGCTTTTAT AAAGATTGTT GATAGTACCC ATACTATTTT TTTCTTCATT 1620

GTATTCAATA TTCAAAGGAT GATTATCTCT GCCAAGTATT TCTTGCTTTA TTTTATTTTG 1680

TTCTTTTATG CTTAGCTTTC TATAACTGGG AATATGTTTT GAGAAAAACT CTGCTTCATT 1740

AAACTTAGTT TTTTCCCATG GAGGATTTCC AATTACAATA TCAAATCCTT CTTGAATATC 1800

TGGAAACTCA ATTCCATAAT GGAAAAATTT ATAGTGGCTA CTTAATTTTC TAATTTTCTC 1860

CATTTTTTCA TTATCTTCAC TAGAAATTTT ATTGCCTAAA ATATTCTCAA TTAAACTAAT 1920

TACAACTGCA ATATCACTAA ATTCTATATT TAAAGATTTG TCAAAAGATA ATGAATAAAG 1980

TTTAATTAAA GAAAATATTA TTCTTAAATT ATCTATATAT TTACTTTCTT CATATTCTTT 2040

GTATATCTTT TTAGATCTTT CTATATCTTC CTTAGTGGTA TCGTTAATAC CTTTAATTTT 2100

TTGATAACCA TCTTCTAAAA TAGTTGTAAT TTCCTTAATT CTTTTTTTAA ATAAAGAAAA 2160

TCCACTTTCA AATTTCTTTT TTGCAATATC AAAAAATTCA TCTTTAGTAT ATCCCAAGAG 2220

AGCATTTCCT GTTTTTATAT GATGCTCAAT AAAGCTTAGT GGCGTTCCAA AAATAAAGGT 2280

ATTAATCCAC AAACTTAGCA TAGTAATTTC AACCGAAATA GGAT 2324

(2) INFORMATION FOR SEQ ID NO: 56:

(i) SEQUENCE CHARACTERISTICS:
(A) LENGTH: 2148 base pairs
(B) TYPE: nucleic acid
(C) STRANDEDNESS: double
(D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 56:

TnAAATCTAG GATACCCCGA cAAATATTAT TTTCCAGTAA TTATGAATAT TTGCTCGTAC 60

GACAATGTAA AGAAATTGCC TTATGACGAG CTTTTAGAGG TCAATAGACT TGCTGAGATT 120

AAATTAGAAA AAGAATTGTA TGAATTAATT TTAAGCAAGT GAGGGCTTAG TGAGCGACAA 180

ATTCACCATT AAATTTAAAG GGATTCTTGA TCATGCTGCA ACAAAAAAGG CCATTGAACA 240

AGATATTTCT AAAATGGAAA AATATCTTAA ACCCAGAAAC TCCAGTTTGG GAAGCACTAA 300

AGATATTGTA AAAAATAATT TGTCGrACAA GAAAAAAGAA CTTAGyArAC AATCTAAATT 360

TGAAAGCTTA AGAGAGCGTG TTGAGAAATA TAGACTTACA CAAACTAAAA ArCTTATAAA 420

ACAGGGCATG GGrTTTGAGA AAGCTAGAAA AGAGGCTTTC AGAAGATCTT TAATGTCTGA 480

TAGAGACAAA AGGCrTCTTG AGTATAAAGA ACTTGCAAAA GAATCAAAAG CAAAAAGTAA 540

AATGTTAGCG GCCTCTCAAG GAAAAGGACT TGTTGCCAAA ATTGCAATAG GTAGTGCCCT 600

ArGGAATATC ATTAGCAACG CTATGAGTAA AGTTGGAGGA GGCCTTTTAG GTTTTGCTAA 660

AAAAGCrGTT GAAGAAGACA CCAAAACAAA AAGAACAAAA CTTCTCAATA GTGCATTTTT 720

TACAGATAAC GAACGAAATA TGATTATGGG AAATAAAGAC AAGAATACTA AGGGAATTCT 780

TGACGGAATG AArGGwTTTG AGCGmGACTT AGAAAAAGAA GArTTcTtAm ATCAAGCAAG 840

TGyCTTTAAG GGTACTyTAA GGGACyTAGA TATGTTAAAT GAAACTAATT TGAAAAACGC 900

aGTAGAATTT GCAGCTATGC TTAAATCCAG TGGTGCTATG AGCAGCGAAG ATGCAGTAAA 960

GGCTGTTAAT AGTGTTCTTG GGGGTGATGG AAGTGAGCTT TTtGATCTAT TAAAGAAGTC 1020

aGGTGTTGGA GACAAATATA TAGAAGATGC CAAAAkGGCy TGGCAArGCG GGGCwsArGT 1080

AGATCTAGAG TCyAGAATTA CCAAGATGAT GGAAATGTTC GAGGATTTTA AATCTTTCGG 1140

CCTTACAAAA AAAGTCAATA ATGCTGAGAG TATTCAAAGT AATTTGGCCT CAGCTGAGCA 1200

AACTCTTCAA AACTTAACCA CTACTGTCTT GGACCCATTA CTTGACCTCA TTAATAAGAT 1260

AACTAATTAC TTTAAAGACT TTGCGTTTGA AACACACATT ATTAATCCCA TAATTAATGG 1320

CATTAAAAGT ATTTTTAATC TTAATTATTT CTTTGCAAAA TTAAAATCGA TGCTACCTGG 1380

ATGGATGGGC GGAGATGAGG GTGCGGCTCT AAAAAAACTA CAAGAAGAAA TTCAAAATCA 1440

AGACAATGCT AACAGCACAC CATAATTTTT ACAAAAGGTA ATTACTTATG ACAAGTAACA 1500

AAAAAATTGC TAACAATGCA GCTAACAAAA TAGATATTAA TAATAAAATT ACTAACAATC 1560

ATGATATTGA AAAGAAAAAA ATCAAGGAAA AAATyAATGA TATTGAAAAG AAAGAAATCA 1620

GGGAGATTAC TCGAATAATA AGAGATGTAA TAACCCAAAT ATTTGCCCTT TTCGGAGCAG 1680

ATAATTTTTT AGTGTTATTT CCTAGAATGG ATCTAAAAGG TTTTGGATAT ATTCCTCAAT 1740

TGTTTTTTAT AAAACCAAAA AATGAACTCA TAACACGCAC TTATAATACT AGTTGTTCTA 1800

AAAGACCAGT TATCAATTAT TATGATAGAA AAGCGGAATA TGTAAGCTAC AATCCGGTAA 1860

TGACTGGTGA ACATATCTCA TTAAACGGkG GAATACTAAC ATCCTTATAT AAGGATATGm 1920

TTTCTTTACT yAAAATGACT GTTTTTGGCA ATACTATGCT ACGTTTTGAC GCGCATCTTG 1980

TAAAAGAACA ACTAGCCAAT AGAATACAAG CACAAGTCCC TTTTAGTATm TATAGTCCAA 2040

CTTTTGGACT TAAAGAATTA GCTGTAATTA CAAGTCTTTC GTTTAAAGAT ACTCCTTTCA 2100

TTGACGAGTT GAAGGTTAGT CTGTCAATAG AAATAGTAAA AACATTCG 2148

(2) INFORMATION FOR SEQ ID NO: 57:

(i) SEQUENCE CHARACTERISTICS:
(A) LENGTH: 2123 base pairs
(B) TYPE: nucleic acid
(C) STRANDEDNESS: double
(D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 57:

ACATATTATT TTGTTTATTT TTACTAATGC TAAACGGCTG TAATTCTAAT GATAATGACA 60

CTTTAAAAAA CAATGCCCAA CAAACAAAAA GCCGGAGAAA ACGTGATTTA ACCCAAAAAG 120

AAGTAACACA AGAAAAACCT AAATCTAAAG AAGAACTACT TAGAGAAAAG CTAAATGATG 180

ATCAAAAAAC ACAACTTGAC TGGTTAAAAA CCGCTTTAAC TGATGCTGGA GAATTTGATA 240

AATTTTTAGA AAACAATGAA GATAAAATAA AATCTGCACT TGATCATATA AAAAGTGAAC 300

TTGATAAATG CAATGGAAAA GAAAATGGGG ATGTTCAGAA AAATACATTT AAACAAGTAG 360

TTCAGGGAGC TCTTAAAGGG GGAATAGATG GCTTTGGTGC AAGCAATGCA ACTACTACAT 420

GCAACGGTTC CTAATAGCTA TTAGCCCCCT ATTTGGGGGC TTTAATCTTG CTATGCTGCA 480

AATATCTCAT CAAACCGTTC GTATTCTTTT AAGATACTAA AGAATATTAA TGGACTAGGG 540

CCATAAATAG GCCTCTTAAG CCCCATAAAC CTTTCAAAAT CTTGTAAATC CTTTAATCTA 600

TTATTTTTCT TGAAATAGTT TTTTATAATC TCGGCCCAAT AGTTTATGCT TTTAAAATCA 660

CTATTTTTCT CAAAATACGA GATTAAATCA GATTCAATTT TCTTGATATC ATCAAAATTA 720

TTGGGGTCTA TATTCTGAAT AAAGCTTATT TTTTGAATAA TTGAGTTTAT ATTATCTTTT 780

GTTGTAGGGC TATTTACCCA GTCTTTTATA TTAGACAAAG CTTCTTTAAA AACACCGTAA 840

TAATAAAGCC TATCTTTTTT TTGTTCTTGC TTAATATCTC TTTTCTGGTT TTGAATAGTG 900

TTATTTTTTT GAATTTGATT AATCTCTTGT TTTTGTTCTC TTTTTTGTTC TTGCTTTTGA 960

TTAACACTAA CTTGCTTGCT AGGCATAGAA TTTTCGTTTT CGTGGTTATT GTAAATAGGA 1020

GCTGCATCAG TATCTATTTC ACTTTCTATA CCAAGAGCTG CAACTAAAGC ATACCTTTTG 1080

ACATAAGTAA TGCCTGAACC AAACATCTGA TACACTGTAT TTGTAACTTT AGACCCATTT 1140

TCATTGTTCC ATTGTAAATT TTCTGTAGGA ATTCGCGTAT CAAAAGAAAA TTCATATCCA 1200

GTACTTGTAC TGTAGAATGT AGTCCTAATA TAATCAACTA TGCCATATTG ACCCTCTATA 1260

GAAATTGGAT ATTGCTCAAT ATCAAGCTCC AAATTGTGCT TTTTAATAAC ATTTTTAATT 1320

TCTCTAACTA TTTCATTGAA ATTTTGATAT TTATATCCAT ATCCTTTAAG ACTTTTGTCA 1380

ATCCCTGGTA AATTCATTTT TAGGGTTTTC ATATCTTTTC GGAAGCTTAT TTTTGCTTGA 1440

ATATTATTTT GTATTTCTTG ATTATTGTTT GAAAGATTTT CCATCTTTTT ACTCCTATGG 1500

TTATTTATAA AAATAAGTAT ATAGCAAAAA CTATTTTTGC CAACTTTTTT ACAAAAAATT 1560

TTACAAAAAA ATAGGGCTTA GCTAAATTCT CTATTATCTA CTAAAGAAAT TAGTTAAGCC 1620

CGTGCTAAAA ATTTTTTTGC AAATTACCAT AGGTAGTCAA AACTGAAAAA TGTTTAAATA 1680

ACTACGCTGT TTGTAGTGTA GCCCAATTTT AAATTAAAAT CAATTTATAT TTTCACTGAA 1740

TTAAAAATTT CTATATTAAT TTAACAAAAT TAATAATTAA AATTTAATAT TTTTTTAGAA 1800

AAGTATTTAC TTTTAAATCA AAATTTTGCA TTATAATAAT TAATTATTAA TTACCATAGG 1860

AGAAAAAACA TGAAAGGTTT TTCAAATACC ACAAAAAATC CCACTTGCCA CAACAAACAC 1920

CAACACAAGT TAATATATCT AGCTTCAACA CTAGATTTtC TAAACAAAAA AGaTAAGAAA 1980

TACACACAAC AAAACATACT CTATTACTAT AATGrrAATC TAAAAAGAAA TGGkCTAGCT 2040

CCCACTACTC TAAGrACmAT GgCAAAATTA TCTTTACAAA TTAGAAAAAG TATTAAAAGT 2100

CACAACTAAT TACTAACCCn AAA 2123

(2) INFORMATION FOR SEQ ID NO: 58:

(i) SEQUENCE CHARACTERISTICS:
(A) LENGTH: 2093 base pairs
(B) TYPE: nucleic acid
(C) STRANDEDNESS: double
(D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 58:

AAAAAAGTGC GCCGTGCGGC GGTGnAACGA CTCAATAGTT ACTAATAATC CATTTCTAAA 60

CTTAGAAATG CAAATTAATA ATATTTTCTA AAGTATTTTT TGTTTCCTCA TAATATGTTT 120

CTTTTACAGA AGGTTCTAAA AGTTCATTTA TAAGAACTTT TACACTATTG TAGTAATGAA 180

TTTTTCCCTT TATGTAAAAA GAATATTCCT TATAAAGCAG TTCTTCTACA TCTTTAAATG 240

TATTTCTATT TTTTAGAAAC TGATTTTCTA TAATCGAAAT ATTAAAGCTT TTATTTCTAA 300

AATCTTTAAT ATCATTAATG GTTTGCATTA AAATACTTAA ACTTTCTACA GAGAATCTTT 360

CTACTTGAAC TGGAATTATT ATATAATCTG TAACATTCAA AGAATTTTTT AAAATAAAAC 420

CCAAGTTGGG TGGGGTGTCA AGTAAGATAT AATCAAAATT ATAATTTGTA ATATTTCTAT 480

TCAAAATATT TTCTAAAAGA AGATCTTTAT AATTTAAAAT TTCTGAATTA AAATTTTCTA 540

AAATAGGATG AGATGGAATT ATAGAAATAA AATCATTAAT TTTATTAATA CACTGTCCAA 600

AATAAACATC TTTTTTTAAT AAGCTGTAAG AATTGCATTT ATCAATGTTG AAAATATATT 660

TACTAAAATA AGAACTTAAA GAATTCTGTG GATCTAAGTC AATCAATAGA ACTTTTTTGC 720

TTAAATCTTT TAATATATAA GAAAAAAGTA TTGTTAATGT GGTTTTACCT ACACCTCCTT 780

TAGGGCTTGC AATTGTTATA ATGTTTGATT CTTTTCTATC CATTTGTTTA TTATTCCTTC 840

CTCTTTTATT TTTTTATTGT AAAATTCGTA AACTGTTTTT TCCATGTTTT TTATATTTTC 900

TAATGTAAAT TTATAGTATT TTGTTTGTTT TTTTTCTTTT CTTAGTAATG TCCATAGGGA 960

CTGAACGTAA CACTTAATAG ATCCTTTTTT AAATACATAT TCTATGTAAT ATGCCTTTTT 1020

TACTACATAT TTTATATTAT TATTGCCTAT GATAAAAAAA GGTTTCTCGA GATTATCCCA 1080

ACCGTATTTT ATGCCTAAAA ATTTTTCATC TTCTTTTATT GGAAATAGAT TGAAAAAATT 1140

CCATCCTCCA GTATCATTAA ATTTTTGGAA GGTTATCCTA AGACCCTTTC TAGTAATTTC 1200

AAATTTGATT AAATGCTTAA ATATTATTGA ATAATATGTT TTTTTATTCT CCTTTTCTTC 1260

TATTTTATAA AAGAAGATTC TTTTTTTTGT TTTCTTTTTA AGATTTCTAA ATCTTTCTGT 1320

AAAGTTGTTC AAATTTTTTT CCTTTTATAA TTAGCTAGTA AATCAAATAG AAATGTTTTA 1380

TTTTGATTTA TTTTCCATTT ATTGATGTAA GTGTCTTTAA ATTCTCCTAC AATTTTGAAG 1440

AAATCTGTTT CGTAGTGAGC ATTTTGTCTC ATGAAGTGAA TTTTATTTTT ATATTTTTTT 1500

AGCTTTTTTA AGTAAAACTC TTTTATATCT TTTGTTTGAT AGTCGTATTT TAAATGCTTG 1560

TTGAAATCTT TGTAATTTAT AAATAACAAT GGGGTTATTA TCAAGGCTGG TGGTGGGCCA 1620

CCAACCGCAA TTAATGCATA TGCCCTCTCT GATTCATTTA CAATCAATTT TGCACCCGAA 1680

GATGAACATC AAGATCAAGC TAAACATCCC AAACAAGAGT ATTCTATTAA TTTGATAAAA 1740

GTTGCAATTT TTGGCAATAG AGGCCTTGAG AAAACAGTAA CACCTGAAGC TGGTGGTTAA 1800

GCCTTGGGCA ATAAAAGGAG TTAAACAAAT GGCAGACACA ACGCAATTAT TAAAAGATTA 1860

TCAAGATAAA CGAgTAAACT TGAAAAGTTT ATGAAAAATC CCCAGTATGA CGCTGGTTTG 1920

CTTAGCAATT CtGTAGAGTT TAGAGATAAA AACATACAAT TTTTTGCCTC TGGAGGCACT 1980

AGAACCAGCA AATTtGACAA aTTAGAAAAT CATCCATTTT CtGGGTATCC ATACmAGCGg 2040

GGaTAAAAAG AGTTATTCAA GAGGAAAAAG CTGATCAAAT TCACTATGGA ACC 2093

(2) INFORMATION FOR SEQ ID NO: 59:

(i) SEQUENCE CHARACTERISTICS:
(A) LENGTH: 2019 base pairs
(B) TYPE: nucleic acid
(C) STRANDEDNESS: double
(D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 59:

GTCTTAGTCA TATTTTCATT AAGTCTTTTT GCTTATTCTT AGCAACCTCT AGCAAACTTT 60

TAGTACTACT CATTGAAGCT TTAGCATCAG CCATAAATTG GTGGTAGCCA GCATAGTAGT 120

AGCTCTCATG CCTTGAAGCA TTGCTATTAC TTAATGCCTC TTCTAAAGCC CGATCTGCTT 180

TTCTTTGTGC ATACTCAAAA TCGTTCTTAG CTCTTTTTAA AGCAGCAATA GCATCATTGC 240

AATGAGTATC AGCAGAAGCA TGATTACTCT TAACCTTAGC AATAGCTTCT TCTAGGCTAG 300

GCAATAAGGC TAAGTTAGCT TTACTAGATC CCACACCTCT TCTAGTCTGC TCTAAATAAG 360

ATTTAGCTGT ACTAAGTAAG CTTTTTATCT TATCAAGGCT TGCTTTTACT TTTGCTAAAT 420

CTTCATTTAT TCTATTGTAC TCTTCTTTAG AAGATTCAAT TTCCGTCATA ATGTTTCTAG 480

CTTCACTAGC TTCATTATCT TGGATATTAA CTTGTAGAGT ATGGTTGCTG CTACTATTAT 540

CATTTTGCAT ATCGGGTTGC ATGTCTGCCA TCAAAGGTTT TTTGCCTGTA TCTTGGTCCA 600

TATAAGTATT ATCTACACTT CTAGGCTTAC GACCACTTCT CTTGCTGTCA TCGTTACTCT 660

TTAGAGTATT AAAAGACCCA TTATTGCTTT TTAGTTTGTC TATAACCTGA TCGGCGTTGT 720

ATGTAATCTT CTTGTATAAT TTACAAGAGA TAACACTGGA AAATAAAATA GTTAGAATAA 780

ATAATTTATT AATTAAATTC ACCTTATTAC TTCTCTTTAG AATCTGATAT TCTTATATTA 840

AAAAATATAT TGCATTTATA ATTTATAAAA TGATTAATAA TGCATAAGCC AAACTATAAA 900

TCTGAAAGAA TTAAAGTAGT GTTTGTAAAG GCAAGATAGG AGTGAGATAG TTAATTTAGG 960

TAAGCTGATA TAAGTTTTTC TAGTAAATAG AGTTTCATAT ATGATTAGAG ATATTAAAAA 1020

AACTAAAGCC TACACTGGAA GTTTTAAAGC TTTAATATAT TTTCTATTTC TTTTCTGCAA 1080

GAATTTCCAT ATTGAAGAAA ACTAATAGCA TTTTCCATAT ATTTAATGCA TTCAGTAGCA 1140

AGATCTATAG CTCTTTTTGC TAAATCTTGT ATATTGCGTT TGTTAAAATT ACTGCCTAAC 1200

CAGTATGAAT TGCCATTTTC ATTGTTTAAT CTTTTATAAA TAGAATCTGC TAGTTTATCT 1260

CGAGCCACAT CTAGCATATC TTTAGCAGAC CCCCAAGAAT TAGTTGCTTG CTCCATAAAA 1320

GAACGCATTT CTTCTAATTT TACCCATAGT GTGGCTTTGA CAGACTTTAT TTTTCTATTT 1380

AAAGGAGATG TGCTTAGATT AGAAAATAAT AAATCTGACA TTTTTTGATG AAATTCTTTA 1440

AACTCAAGGA GTGTATTGTT TAAGTTATTT TCAGAATTAG CACGATTAAT AAACTGCTCT 1500

GCATTTTTAA TTTCTGAAAG AGATGTTTGC TCATACTCTT CTTGTGCAAA TTCATCTAAT 1560

TTAGCTTTAA TTTCTTCTTC TGAAGGAATA GCAGGAGTTT TACTTGTTGT AGTAAGGCTA 1620

TTTTTAAAAT TACTTTGTTC AATTGGATCA TGTTGTGGAG TAGCTGATTT ATTTTCTTCC 1680

ATCTTTACAT TATTTTGAGC ATGCGAGTCA TTTTGCATTA TTGCTGGACT TGAATTTGGT 1740

AGGGATTCTT TATTATTTGC ACCAGGATCA TCTGACATAT GTTGATTAGA ACCTACTTGT 1800

AAGATTGGAT CATAAGTTTT TTTATTTGAT GATGTAACGT CTTTAGGAGA TTCAATTTCC 1860

TTACTATTTT TATCTACTGA AGTATTAGTA GTATTTTTAT TTTCTTGATG ATCTTTTAAT 1920

GCTAATGCAA ACTGTTCTAT AGATTTTTTA GATAGAAAAC CAGAACAACT TTCAAATAAA 1980

AAAAGAGAAC TAAATACTAA ATTAGGTATA ATAAAAATT 2019

(2) INFORMATION FOR SEQ ID NO: 60:

(i) SEQUENCE CHARACTERISTICS:
(A) LENGTH: 1907 base pairs
(B) TYPE: nucleic acid
(C) STRANDEDNESS: double
(D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 60:

AGAAAAACAA AATCTATTGC TTCTTCACTA AATCCTATCT TTAAAAATTC ATTTTTTATA 60

CTTTCTATAT TGTATGTTCT GTAAGTCAAA CTATTCATAA AATTTCCATA TTATCCTTTT 120

AATTCTTTAT ATTCTTTCAT AAGTTTTTTA ATTATTTCTT TTCCATCACT AAATAATTTA 180

TCTAACATAA ATCCTGTAAA TTTAGCATTA CTTTTATAAA AATCATAGCT TTCTTGTCTT 240

TTAAGCTGAA ATCTTAAGGG CTTTATCGGG TTTTGTTTTG ATTTTTTTAT TGTTTTGCTC 300

TCTTTATTTC TTAATACAAT TAACGTTTCC AGTATACCAT TTTTTATTAG AAATTCTTCT 360

TGTATAATCC CATCTTCTAT TGCATTAGCA ATTTTTAAGT AATTATAGAC CTGTGCTCTT 420

GCAAGTCTAT AATCTTTAGA AAAAGCTTCA AAACTTTTAT AACCATCAAA CCTATAATAA 480

TGATTGTCTT TAATTTCTTT TAAAATTTTT AAAGTTTCTA ACTTACAATA GATTTCCTTT 540

TTAGAATTAA TTTTTAACTT TTCTTTTAAA GAATTATAAT GATTTAATAC ACTATCAGTA 600

ATAATATAAT TTTCATTATC ACTTAAATCT CTTTTATTAA CCTTAATATC CAATTTAAAC 660

TCCTTTTACA TTAAACTGTC TAATTATTAG ACTTTATATT TTTTTTAAAA AAATTTCTAA 720

AATATTTTCA TATTCTTTTA TATAATCTTT ATTTAAATCA AAATTATTAT TTTCTGCTAT 780

TCGTCTATTT AAGTCTTCTC TTTCAGATAT TGTTCCTAAA AATCTATCTT TTGTTTTTAA 840

TATTTCAAAT AATGTTTTAT GwGTTCTAkT TwwwtTaAAT CTTGTTATTA TCAAAAATAT 900

AGGTAAAAAT AAATTTAATT TTCTTACAAA GAAATTAAAT AAATCTAAAC TTTCCACTGC 960

CCACTTTTCA GCCGTCATTG GAATTATTAC ATAGTCACTA CATAAAAGAG CATTTTTTAA 1020

CGTAACATCT AAACTGGGAT TTGTATCGAT TACTATATAG TCATATTTAT AATACAAAGT 1080

TCCCAGGCTG GTTTTTAACA AAAAATCTTT ATGTTCGATT TTATCTTCAC TAAAATTATG 1140

TAGCGTAAGA TAGCTAGGTA TAAGATCAAG ATTATTATCT ACATTTATAA TGGTACTATC 1200

GATATCTACA TTTTCTTTCA AAATCTCATA AATATTAAAT TTGGTAAAAT TAATACCTAG 1260

TTTTTCTATT TTTTCGTAAA AATAACTAGT AATAGATGCT TGAGTATCCA TATCAATTAA 1320

AAGAACTTTA TTATTTTTTG ATAATAAAGT AGCCAAAATT ATCGCACTTG TGCTTTTACC 1380

TACACCGCCC TTAATTGACG CTATTGTTAT TATTTTAGGT TTTTTATTAT CCATTTTATT 1440

AGTGGTCCTT GTTCCGGGTA TTTCTTCCCA TAAAATTTAT ATACTTGTTG TTCTAAATCT 1500

GTAAACATAC TAAATAACAC TTTGTTGTAG TGATTGTTTG TTCTTTTTTT ATCTAATAAA 1560

CGATATAATC CCTTGAAATA GCAAAAAACA CTTCCTGCTT TAAATCTAAA TTCCATATAA 1620

TATGCCCTTG CTAATGCATA TGCTTTTCTA GCCCCGTTTA TTTGATACTT TATTAACGGC 1680

TTTTTAATCG GCTTTCTATA GCCATAAAAA ATACCAATAA ATTTATCTCC TTCTTTAATT 1740

GGGTACAAAT GAGTTTCTTC AACAATTCTT TCTCCATTAA ATAAGGCCCT CAATGATAAT 1800

CTAAATTCGT GTTTTTTCTC ATAAACTCCA AATTTATAAA TATCCATCAT TATTTTTGTA 1860

TGGTACATTG CTTTACCATT TTCTTTTTCA ATTAAAATAA AGCGTTC 1907

(2) INFORMATION FOR SEQ ID NO: 61:

(i) SEQUENCE CHARACTERISTICS:
(A) LENGTH: 1907 base pairs
(B) TYPE: nucleic acid
(C) STRANDEDNESS: double
(D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 61:

GAAAAGATAT ACGTAGAAAT AGAAAGACGT ATTGAAAACC ACAATTTTTT GTTTTACAAA 60

GATGAATCTT TAGTACAACT ACAAGACGCA CTTTCTAGTG CAACAACTTC TTTAAGTGCA 120

CTTACTCAAG GCAATAATGA TAGAGGAAGT GGCATTTTAT CTTCTTTTTT AAGAAAACAA 180

AATTCAAACA ATCATAGTAA AGATATTTCT AATTTACGTA GTCTTAATGA CTCATTGGCA 240

CAGGAACTTG CTAGGTTAAA AAGCAATCTA AATAATGAGG GAATGTTTTA TACAGCTACT 300

CCGAGTGCTA GTTTAGAGGT TATTAAATAC GATCTTAGCT ATTTAAAGGA GGCTTTAGCA 360

TTAATTAAGG CAAAAATTGG TGCAGATACT AAAGAACCCC TAACTAGAAG TTTTAATGAG 420

CAGGCTAAAG GACTAGGGAA TGATGGTAAA GGAGATAGGA GCAATTATTA CGATTTTCTA 480

AAAGGTGTAC AAGAACAAGT TGAGAACTCT TGTAATTTAA AACTTACAAA GTATTTTGGA 540

CTTGATATGA AGTTTAATTC GCTGATTATG TTAAGTGAAG AACAAAAAGT GGAAAGAGAT 600

ATAAAGCTAA TTGAGCTTaC AGTAAATATA ACCAGCTTAT ACAAAGTAGC TCCTTgATAA 660

TGAGGAGTTA GCGATTTaAA AGAGAAATaT TCTCATTTtG AGAAAAGGAG TTAAAAAGTG 720

ACTGAGAAAG AAGAAAAAGA AGACCTGCAG GCACmAGATA AAGArGAGCa GCAAaTTAaG 780

GCtGATACTA AAGTTATAAG TGCGCAGGAA TTTGAAGAGT ACATGCGTTT TAAAGAGCAG 840

GCAAATAGTA AATCTAAAGA GACAAGTCGA GATTTAAGTA TAAATGAACG AATAACAAAA 900

GAACTTGCAG AAGTTGAAGA GCGGGAGCGT ATTGAAAAGC AATTGTTACT AGAGGCTGAG 960

CGAATTAATG AAATTGATAC ACTTGCAAAA GCACATCTTA GCAATCATTT TAACAAAGAG 1020

GTGCTACTTG CAAAAGGATA TACATTAAAA GACATTATGC AAGCACAACG TAGAGAACTT 1080

GTACGCAAGT TCGTTCCAAT TGAGCAAATT AAAGCTATTG CCAAAGTATC AGACATAAGT 1140

CATATmGATG GrGAGATATT AGAGCAACTT GTTTCTTTAG CAAAAGTGAA TATTAAATTA 1200

AGAAAAAATG CGAGTAGCAA TTCTTCTTCT GTTGACTCTA TTAAGGGGAA TATTGCTATT 1260

AAATCAGAAG AAAGAGCAAG TTTGCTTAAT TCTAATTTTG TACCTATTAA TTTCACAGAA 1320

TTTGTACAAG CGATAAGTAA TACTTACAAG CAAAGACGAA TTCAATTTTA TGAAAATCTA 1380

AAAAGACATA AAAGAACAAG TATTGCTTAA AGGAGTTTTT AATGAGCGAT GGTATTACAA 1440

AAATAAAAGA AGAGTTTGAT AAAAAAGTTG CAGAAATTAA AGCATTAATG AAAAATCCTC 1500

AGCAAGATTC TGGTTTGCTT AGTAATTCTT TAGATTTTAG AGATAAAAAT CTAATTTACT 1560

CCAATTCGGA TGGAGTTTTT ACTAGTAGTA AAGACAAAAT AGAAAATTAT CCTGCTAAAG 1620

GGTATCCATA CAAGCGTGGA GTCAAGCTTA GTTTTAGTGC AGATGGTACA ACAGAACTAG 1680

AAGTTGAGGC TGGTGGTGGG GATGACTTGT ACGGAATATG CACTGATATA nATGAGTTTA 1740

CTGGCATGGC AACTGTAGTT CCAATTACAA ATAACTTCAC AGGGTATTTA ACATTTAAnA 1800

AAAATGGACA AAATGGTGTG AATCCGGGTG ATAAGCTGCA TTTTAATGCA CnAGGAGAGC 1860

TTGAAAAGAA TGGGGGAAAT GATAAATCTG TTAATGCTAT AGCnCTT 1907

(2) INFORMATION FOR SEQ ID NO: 62:

(i) SEQUENCE CHARACTERISTICS:
(A) LENGTH: 1902 base pairs
(B) TYPE: nucleic acid
(C) STRANDEDNESS: double
(D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 62:

CGGGCATTTA AGCTCTTTTC TACATTGTCT ATTTTGATAT TCAAACCATC TATTTTTAAA 60

TTTAAATTCT TTTCCACATT GTCTATTTTG GCATCTAAAT TAGATATGTC TTTTTGCAAA 120

TTCTTCTCTA TATCAATTAT TTTCTCTTTT AAAAATTCAA AGTTGTAATT ATCATTATGC 180

AGAAAAACAA AATCTATTGC TTCCTtGCTA AACCCTATAT TTAAAAATTC GTTTTTTATA 240

CTTTCTATGT TATATGTTTT GTATGCTAAA TTGTTCATAG ATTATCCTTT TAATTGTTTA 300

TACATTTTTA AAAGTTTACT AATCAAATCT TTTTGATTTT CAAAAATCTC TTGCATCATA 360

AAACTTGTAA ATTTAGCATT GCTTTTGTAA AAATCATAAC TTTCTTGAGT TTTAAGTTGA 420

AATCTCAATG GTTTTATTGA GTTTTGTTTA GATTTTTTCA ATACTGGACT TTCTTTATCT 480

TTCAATACAC TTAATATTAA TCTAAATCCA TTATCTAATA CATATTGTTC CTCAATAACT 540

CCTGCTTCTA TTGCATTGGC AATTTTTAAA TAGTTATACG CTTGAGTTTT TGCAACATCA 600

TAATCCTTTA TAAAAGCATC GAAACTTTTG TATCCATCAA GTTTATAGTA TTCATTATCT 660

TTAATTTCTT TTAAGATTTT CATACATTCT ACTCTATTAG AAACTCCTTC TCTAAGGTTT 720

ACATACAATT TCTTTTTCAA AGTATTATAA CGATCTGTTT CAACACCATT TTTACTAACA 780

TTAGAAGAAT CTACAAGTAA TGCATTCCCC TCAGAATCAA TATCCCTTTT ATTGATTATT 840

AATTTTGTAT TATTTTTCAT AACAAGCCTC CTTAATTATA AGTTCAACGC GTCGAACTTA 900

TAATTATAAT TATTTTAATT TTGCATAAAA ATTCATTAAT GAATTTTTAT ACTCTTTTAT 960

ATAATCCATT TGAAAATCAA AAGAAGAATT ACTAGCAATT CTTCTATTTA AATCTTCTCT 1020

TTCTGATATC ATTCCTAAAA AATTTTCTTT GGAGTTCAGC ATTTCCAACA ATTGCTTATG 1080

TGTATTATTT TTTTTAAATC TCGTTATTAT AAAATAAGTA GGCAATTCTA CACCTATTTT 1140

TTCCATAAAA AATTTCAAAA GGTCAAAACT TTCAATTGTC CATTTTTCTG CTGTCAAGGG 1200

GACAATTACA TTGTTACAAC AAACTAAAGC ATTAGTTAAA GTAAAATCCA AACTTGGGGG 1260

AGTATCAATT ATAATAAAAT TATATCCAAC ATCTATATGT TTAAGCTCTT TTTTTAATCT 1320

AAATTCATCA AAAGTGTGCT TATAACCAAA AGCATTTATA CTATGTAAAG TCAAATAACT 1380

AGGTATTAAA TCTAAATTAT TCGCTACATT AACGATTGAT CGATTAATAT CTAATTTTTC 1440

TATTAAAACT TCATATATAT TATTTTTTCT TAAATCTATA CTGGATTTTT GTATATCATC 1500

ATAATAATAA CTAGTGGTGG ATGCTTGAGT ATCTATATCT ATTAATAATA CCTTATATTT 1560

TTGAGCCAAT AAGGTTGCAA ATATAATTGC ACTTGTGCTT TTACCAACAC CGCCCTTGAT 1620

TGACGCTATT GTTATTATTT TAGGTTTTTT ATTATCCATT TTATTAACGG TCCTTGTTCC 1680

GGGTATTTTT TCCCATAAAA TTTATACACT TGTTGTTCTA AATCCGTAAA CATACTAAAT 1740

AAAACTTTGT TGTAATGATT ATTTGTTCTT TTTTTATCTA ATAATCGATA TAATCCCTTG 1800

AAATAGCAAA AGACACTTCC GGCTTTAAAT CTAAATTCCA TATAATATGC CCTTGCTAAT 1860

GCATATGCnT TTCTAGCCCC GnTATTTGAA CTTATTAATG GC 1902

(2) INFORMATION FOR SEQ ID NO: 63:

(i) SEQUENCE CHARACTERISTICS:
(A) LENGTH: 1761 base pairs
(B) TYPE: nucleic acid
(C) STRANDEDNESS: double
(D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 63:

AACnGGCCCC GGAAGTTAAA GCCTTGTGGG nCCCATGCtC TAGTGGATGA CCGTTCTTTT 60

AGAGCCTAAA AAGCTATCAT GGGATGAAAC AAGAAGCTAT TTCTATAATC TTTGATTTAG 120

AAATAGCAGT TCACATAAAG ACGCTTGATA TTAATTTAAA TTTAATCTAT AATATAAATG 180

GCGTAGTATG AATGTAAAAT AATTTACGTT TGAGCTGCCT TATGGAATCA TTTACAATTG 240

AAGGGCTTAA AGAAGTTCTT TAGCAAGAGA GTTTCTTTAA GCCCTAATAA TATTTGAACA 300

ATCTTTTTCT AGGTAAATTG ATCTTCAATA GGATTTTTTA AACGACAGTT GGAATCCGTT 360

TATTCCAATG TCAAAATTGG GTTCAACCCC CGCAAGTGCA ATGCCGAGCC TTTTTTTAAG 420

GTCTGCGTTG TATCTATTAG CAAATTTAAA TGGAATAATA ATTCCAGTTA TGTAGGATGC 480

TACAATTGTG AGCCCTCCTA TTCCTGATAA TACTCCTCCG GTTATTACTG TTGCTGTGCT 540

TTCTGTAACG CCTCCAATAC CTCCTACGAT CATGTGTCCA GCCATTATAA GTATTCCTTC 600

CAAGCACTTG AGAGCCAAGT AGTGCACCAC CACCAATATA ATCTCCTTGA ACAAAAGATC 660

CTATCCCTAA AGACAAAAAG ATATTCAAAA GTAATGGTGC TAGTATGGTT GCTTTTTCGC 720

TTTCATATTT CATTACAGTC GCGATATCTC CACTCCAACA CCTTTTTCAA GTTTATCTTG 780

TGCAAAGATT TGCATTGTTA AACTAAAAAT TAATATTAAT GTGAAAATTT TTTTCATATT 840

AATATTACCT CCTAATAATT AAGTTTTGAT AAACAAATGT TAGCACAATT TTTAGATTTT 900

ATTTATGGAG TTGAATCTTT TCTTAAAGAT ATTGTTTGAA TTTCTTTGCT GTTCTAAGCA 960

GATTTTAATG TAAAGTTTTC ATTAAACTCT TTTAAATTTG AAATTATGCT ATCTATTTTT 1020

TTTGTATGCC AATTGTATAA TGAATTGTTT TGATGACTTT TTCGAGATGT TCAACTTCTT 1080

TTGAGGGCAT TTCTCTGCCT TTTCTATAAG TTAAGTAACT TTTTAGAGTT TGCCAACTCC 1140

CAATAGTATA GTTATACACC TCTTTGGAAA CATTAATAAA TCTAGAAGTT TGGTTATAGT 1200

AAAGTTCATT TGTTTCTTCT TTATAAAAAA CTTTTTCTAT GATTGAGTTT TGTTTTAGGG 1260

TTTCATCTAA AAATGAAAAA CATTTTCCAA TGTTGATATT TATTGTTGGA ACAACTTTTA 1320

CTAAATGAGA ATTAATGAGA TCTGTTCCAA GTTTGCCAAG TGTTACAAAT ATGTCAGAAT 1380

TATCTACGAA AATAATTTTA GGAAAGTCTA TTTGTAGGTG TTCGTAGAAT CTATCTCGAT 1440

AGATATTTGA ATAAAGAATT GCGTAAATAT AGCCAAGTAT TTCTTCTGCA GTAAATTTTT 1500

TATTGTATTT AACATCAAGA AAATGTCTAA ATTTATTTTT AAAGTTTTCT TTTTTAACTC 1560

TCTCAGGTGT TTCGCTATCT TCTTGTATAT AAATTGGGAA AACATATCCT AGTGGAATAA 1620

TGCTTAATTC TGATATTTTA GAAGTAACGA AAGCATGAGA AAAACGATCA GTTTTGGATA 1680

GTCTTGTTGT TATTAGTGCT ATATTATTTT GAATTTCTAA GATATGTTTC ATTATTTTAT 1740

AACCAGGCCT AATTATGACC C 1761

(2) INFORMATION FOR SEQ ID NO: 64:

(i) SEQUENCE CHARACTERISTICS:
(A) LENGTH: 1717 base pairs

(B) TYPE: nucleic acid
(C) STRANDEDNESS: double
(D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 64:

GCATCATTTC TGAGACTGTT GTTTTTGTAG ATAAAAATTT TCCCAAATAA TATTTAAGCA 60

ATACAAGATT TAGTAAAGCG TATATTTTTA TTTTTTGATT TAATTTCAGT TATTTTTAGG 120

ATTTTTATTA TTTTAATCAT ATTTTCTTTA TCAATATTTA ATGTTAATAA AATTGAAATA 180

ATTTCTTTAC ATAAAAAGTC ACATTTATTG AAATGCTTTA TTACTTGATA CTTTTCTATT 240

TCGTTAATTT TTCTTTCTTC TTTTATATTA TTATTACAAT TCTCCAATTG TACACTACCC 300

ATTTTTGTAT CAGAATTTTT ATTAAAATAG TTGGCAACTC TATTTTGAAA TCTTTTTTCT 360

TTTTTTTCTT TAAAGTGTTG GTTTATCTTA TGGTAACAAT CTTTTTTAGG ATAATTAAGC 420

TTATAATAAA TTTCTGTACC CGAATTTACC CCCATATGTT GATAGTAATT TGTTGTGACT 480

TTTATTTCTT TTTGTAGTCT ATAAATATAC TTTTGCATAG TTCTTAGTGT AGAAATAGTT 540

TGCCCGTTTT TTATAGATTT TCATTGAAAT AATATAATAT GGTTTTTGGG GTATATTTTA 600

GATTTTTGGT ATTTAAATAG CTTGTAGAAA TAAGAACTAT CAATTTATAT TTATATTGGT 660

TATAATTTAG AATATTACTA GATTAATATA TCTAGACTTT ATTTTCTATT TAATATACAA 720

TTAATTAGGA AGCATTATGT GCTCCAAATG GATGATAACC AGATAAAGGG CTTTAAGTGG 780

CTTAAGGAAG ATAAGTTACT TAAAGCCCTT ATCGCATTTA TACTAATTTC CCTAATTTAC 840

GTTTTATTTT GTTTAGAGCG TTATATTATT AATTTTTAAT CATAAAATGG AGGATTAGTT 900

TTGTAATTGT ATATATTTTT AGCTTTAATT GTTCTTTATT AGGCCTTTAA TACTCTAAAG 960

TATTATGCTA TTTCGCAAAA ATAATTTTCT GGATTATATT AGCTTACATA TTAGGATGAG 1020

AATTATAAAT TTTAGTGCAA CAACCTTCTA CAGGGTAAAG GAGTGTATAA AGGCCTCTAA 1080

AGCTTTCATT TTCTTTATTT GTGGAACATT TGAAGATTAC TTTTGGATTT TTTCTAATAT 1140

TTCAAGATAT TCTAACAGGG TTTTAGAATC AACTTCATGT TGATTAGTTT TAGCTAGAGC 1200

CTTATTAAAC TTGCTTCTTA TTAAGGAATA AGCATAATTT TTATTTGCAC TATGGAATAA 1260

TTTTTTATTC ATATGATCTA GTGTTTTGAC AAGAATGATT AACTTATTCT TAGCAAATTA 1320

AATTTGCTTT GTTATTCTTG CCAATGTATT GTTAATTACT CCATCCATAA TGAATTAGCC 1380

CCCTATTATA TTAAATTTAT ATTATAAATA TAGCACAGTA TTTTTTTAAA TTTTTTTAGC 1440

GTAAAACAAT ATATTTCTAA AGTTTTGCAT AAGCTTATTT TATAATGTAT TTATAATTAA 1500

GTAGTAGTGA TTTTTTGTAA AAATTGTTTT AATTCTTCAT AGAATAATAC GTAGTAGTCT 1560

TTTATTTTTA ATAAAACTCT TGCTAAGGAG CAATTAACTT TATAATAAAT TATAGATTTA 1620

GGATTTTTCT TAAAATAAAG TTTGACTATG ACGGTTATTT CTTTTTCTAA TTTGCGTTTT 1680

ATTTATATAT TTTTTAGTTT TAATTGTTCT TTATTAG 1717

(2) INFORMATION FOR SEQ ID NO: 65:

(i) SEQUENCE CHARACTERISTICS:
(A) LENGTH: 1566 base pairs
(B) TYPE: nucleic acid
(C) STRANDEDNESS: double
(D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 65:

AGnCACGTAA AATAACTTCA GTTCATATTA TCATATAATA AATAAAACAT TAAGTACAAT 60

AACCTAACAT TTAAAAAGGA TGTACATTTT AATACAGAAA CTGAAGCTGT TCAACTACAA 120

TTATTGCACT TGAAATTTTT TATATTTAAA TAATAATACA AATAATTATA TTAACAAATA 180

TCAATTAAAT TTATTTTTAC ATCATATAAT AGTGCTATAT ATTGTATAAT ATGCTATATA 240

CTTGAAACTA AAGGGGGGGC ATATTAGTTA AGATAATATT CTTATATTTT TTATTAAGGA 300

GACTAATATG AAAAAAATAT CAAGTGCAAT TTTTACAATA ACTTTTCTTG TTCTTATCAA 360

CTGTAAAAGC GATACTAGAA AAGCTATTAA TTCAATACAA ATCCAAAAAT TTACTTCCTT 420

TGATGGGTTG ATTGATGGCT TTCTACGCCT TAATTCAAAT CCCAAAAAAT CTGAGAGGTA 480

AAAGATTGTT TTAACAGCAT GGCTAAAACA TTAAATAAGG CCAAAGACAA ACTTGCTAAA 540

TTCATTAGTG AAAAAGGTGG CAAGACAACC GAAGGAAAGA ATACTGATAC TGCTAAAGAA 600

GATAATAGCA CAGTAAACCC TATTGATGAT GAAATAAGTA AAATTAACGA TATGATGGGA 660

AAAATGATAG ATGCTGCTAA TACCATTGTT GAAAATGTAG CCGAAACCGT AACTGAAGCT 720

aTGGGAGAAG TTGTCGAGGT TAAGAGTATT GGTAATGTAG CAACCAAAGC CGATGTAAAA 780

AGTGTTGTTG AGATTGCTAA AGGAATAAAG AAGATTATTG AAGCTGCTGG TATTGCCGAT 840

AAATTAAAAG CTGAAGCCGA TAAATCTACA AAGCCAATCA GCGAAGAAAG TAACAACAAG 900

TAAGCGGGCA AGATGTTCTC TGGGAAGCAG GGTGATCAAG GTGGTCGAGT TTTCGATGAA 960

GTCATTCCAC CTGAGATTGG AAGAGGAGCT AATCCATTTG ATATTAAAAA GGCTACTAAA 1020

GCTATTGAAA GTGTTAGTGG AGAGCAGATA TTAGGATCTA TTGTTGTGCT GCTACTAAAA 1080

CCGTTAAAAG TGGTGGTGAG GAGCCAAAGG GGAAGAATGC GGATGAAGCT ACAAATCCGA 1140

TTGAAGCTGC CATTGGAGGA AATGACGATT CGGATGCTAC TGCATTCAAG GGGAATATGG 1200

AAAAAGATAA TCAGATTGCT GCTGCTATTG TTTTGAGAGG AATGGCTAAG AACGGGGAAT 1260

TTGCTGTGAA AATGGGTCGA AAACCAAGTG GTGATGGTGA TAATATTAGA GTTCTTGTTA 1320

ATAATGCTGC TAATAAAACT GTTGATGCTT TATCTAAGTT AGCACTAGAA GCTATTAATG 1380

AAAGCTTAAC AAAAATAGCC AAGACTATAC ATTGAAAAAA TAAAGTAAGA ATCAGCATTT 1440

TTAATAAATA ATATTATTTA TTAAAAAATG CTGATTCTTA CTCAACATCT TACGTCAGTA 1500

GTTTACTAAA CTGCATAATC ATTACATATA CACCAACATA TCTAAATTTG CAAACAATCA 1560

TCTTAG 1566

(2) INFORMATION FOR SEQ ID NO: 66:

(i) SEQUENCE CHARACTERISTICS:
(A) LENGTH: 1552 base pairs
(B) TYPE: nucleic acid
(C) STRANDEDNESS: double
(D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 66:

CTGTTTTGTA AACCAAAAGT GGATTATAAT AATTGGGCCT ACTAGCTTGA ATTCTAGAGT 60

CAAGCAAACT TACACTAATT GTATCTTGCG GCAATTTTGT ATTCCTCCTT TAAAATTTCA 120

ATTGCTTTTA CACTAGCATT AAATGCTATA GATGCACTGT ATGCATGGTT GCTATATTTT 180

GTGCCTAAAT TAATAAGCCC AACTGTTTGC ATATTAGATA TTGGATAAAT GTAAAAGTTA 240

ACTTTACTAA TATATTCGGG TTGGGATTGA CTTTCTAAAG TATACTTATG GGCTTTATTG 300

TGTAGAAAAT TGCTAAGCAT ACCATAAAGC ATTAACATAC GTGAATTAGC GTCAAAATCT 360

TGAGCGTTTA ACACTATAGC AATAATATAT ATTTGAAAAT TTAAACTGAA TTCCAAAGCA 420

TTTTCATAAA ATGCACCGGC TCTAGAATTA TGATCAAATA GATTTTCTGT ACCATCAAAT 480

TTCAATGCTA TTATATTTGA GCTAGCAGCT GTGATTTTTG AAAGGTACGG GTGATTGTAA 540

GTATTTATGA TATCGCACTC AAAATTATTT TCAGTTGCAT ACGCCTTAAA CCCTTTGAAT 600

ATTTGAGTTA AATGGTTTAA AACCATATCT AAAGTAAAAA TCATTCAAGT GTTACCTTAT 660

AAGTAATCTC GGATAACATT TTGGCTGTAT CAACAAGTGG AATTGCTGCG GTGTTACTAC 720

CCTTTTTAAA CTTACTTTTG ATTGTATTAG CCTTTAAGGC TGGAGAGACT TGTGCTGATA 780

ATAGATAATT TCCATAGTAC CTTATAAAAG CTTGTCCAAT AGCCTCCATT CCCGATTTGG 840

GGTCAAGATT AAACTTAGAA TTTATATAAC TATTATTGAT ATATTCTCTA AATTCAGAAC 900

TACCAGCAAT TTTGGTTAAA TGTTTTCTTG CTGGTAAATT GCTACCCCCT TTTTCATGCA 960

TTCTAGCAAT CCCTGCACGA CCACCAAACC ACCCAATTTC CAATTCCATT TTAAACTCTA 1020

GTTTGTCCAT ATAAACTCCT TTAAAACCAA AGTAAAATAT CCGATTGAAG AGTCAATACT 1080

AAATATTTCA AAGTAAATTA AATCCGAAAT TGATATGCGG TCTTTTAGTT CATAGTTAAG 1140

GTCTTGATAT GTGTAAAGTT TGGAATATCC TTGAATATCA GACATATCAG AGTCATAAAG 1200

CACTGCAAGT TCTTGTGGCC TTATGTCAAT AATAACTCCT GCAAATTCAG TGTACTTATT 1260

TTTATCAAAA ACTCTCTGAT AAGAAGAATC GTTTTCAAGT TTAACAACAG TGCCTTTATA 1320

AAACCTTAAA GGTTGAGGAT CCTTAAATAC GTTGATCATG cGGAAAGACA TATCTGAAAG 1380

TCTTTTTCTA ACACCATTCA TTAGACAACC CCCACACAAG ATGGCGTTGA AGTTTCTCTT 1440

TTTAGTTTTT CTAAAAAgCa TCAAGTTGTG AACAAAAATT CyTGkTTGAG CCACAACCCC 1500

CCTCGsCGGC TTCTTCGGCT CCACTGGCTA CTAGGnTAAT AATCAAGTTC CA 1552

(2) INFORMATION FOR SEQ ID NO: 67:

(i) SEQUENCE CHARACTERISTICS:
(A) LENGTH: 1484 base pairs
(B) TYPE: nucleic acid
(C) STRANDEDNESS: double
(D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 67:

AGTCATTACC GGATTGTAGC TTACATATTC CGCTTTTCTA TCATAATAAT TGATAACTGG 60

TCTTTTAGAA CAACTAGTAT TATAAGTGCG TGTTATGAGT TCATTTTTTG GTTTTATAAA 120

AAACAATTGA GGAATATATC CAAAACCTTT TAGATCCATT CTAGGAAATA ACACTAAAAA 180

ATTATCTGCT CCAAAAAGAG CAAATATTTG GGTTATTACA TCTCTTATTA TTCGAGTAAT 240

TTCCCTGATT TCTTTCTTTT CAATATCATT AATTTTTTCC TTGATTTTTT TCTTTTCAAT 300

ATCATGATTG TTAGTAATTT TATTATTAAT ATCTATTTTG TTAGCTGCAT TGTTAGCAAT 360

TTTTTTGTTA CTTGTCATAA GTAATTACCT TTTACCAAAA TTATGGAGTG TTGTTAGCAT 420

TGTCTTGATT CTCAGCCTGT TCTTGCAGTT TTTTTAAAGc TTCGCCGCCA TCTCCGCCGA 480

ACATTGTAGG TAGAACCGAT TTTAATTTGG CAAAGAAATA ATTAAGATTA AAAATACTTT 540

TAATGCCATT AATTATGGGA TTGATAATAT CTTTGGTAAA ATCAAAATTT TTAATTTTAG 600

CGGTGATCCA GTTAATGATA TTAAGTAATG GGTCCAAAAC AGTAGTGGTC AAATTTGAAA 660

GAGTTTGCTC AGCTGAGGCC AAATTACTTT GAATACTCTC AGCATTATTG ACTTTTTTTG 720

TAAGGCCGAA AGATTTAAAA TCCTCGAACA TTTCCATCAT CTTGGTAATT CTGGAATCTA 780

GATCTACCTC AGCCCCGCCT TGCCAAGCCC TTTTGGCATC TTCTATATAT TTGTCTCCAA 840

CACCTGACTT CTTTAATAGA TTAAAAAGCT CACTTCCATC ACCCCCAAGA ACACTATTAA 900

CAGCCTTTAC TGCATCTTCG CTGCTCATAG CACCACTGGA TTTAAGCATA GCTGCAAATT 960

CTAyGCGTTT TTCAAATTAG TTTCATTTAA CATATCTArG TCCCTTArAG TACCCTTAAA 1020

GrCACTTGCT TGATTTAAGA ATTCTTCTTT TTCTAGGTCG CGCTCAAATC CCTTCATTCC 1080

GCCAATAATC TTTAAAAGAC TCTCTTTCTC TTTTGGATCA CCATAAAACG CTTTATTGAG 1140

AAGTTGTGTT CTTTTTGTTT TGGTGTCTTC TTCAACCGAT TTTTTAGCAA AACCTAAAAG 1200

GCCTCCTCCA ACTTTACTCA TAGCGTTGCT AATGATATTC CCTAGGGCAC TACCTATAGC 1260

AATTTTGGCA ACAAGTCCTT TTCCTTGAGA GGCCGCTAAC ATTTTACTTT TTGCTTTTGA 1320

TTCTTTTGCA AGTTCTTTAT ACTCAAGACG CCTTTTGTCT CTATCAGACA TTAAAGATCT 1380

TCTGAAAGCC TCTTTTCTAG CTTTCTCAAA CCCCATGCCC TGTTTTATAA GTTTTTTAGT 1440

TTGTGTAAGT CTATATTTCT CAACACGCTG GGTACCGAGC TCnA 1484

(2) INFORMATION FOR SEQ ID NO: 68:

(i) SEQUENCE CHARACTERISTICS:
(A) LENGTH: 1452 base pairs
(B) TYPE: nucleic acid
(C) STRANDEDNESS: double
(D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 68:

CTGCTTATTA ATTCCAATAA GTGCTTATAA GTGTTGTTTT TCTTAAAATT AGTTACCATT 60

GGGAAAATCG GTATTTTCAA TTTTAATCTT TTTAGAGCAA ATTCTAATAA TTGCATGCTT 120

TCTACCGACC ATTTTTGAGC AGTCATTGGA ATTATTATAT AATTACTTAC AACTAATACG 180

TTTGTTAAAA TAATTCCCAA ACTAGGACTA GTATCTATTA TTATGTAGTC GTATTTATGT 240

TTTAATAATT TTAAACTATC TTTTAATCTT GTTTCTTTAA ACGGGATGTT ATCGTCATAA 300

AAAAGGTATA AATATATATA ACTGGGCAAT ATGTATAAAT TATTGTTTAA TCTAAAGGTG 360

GAAGAATTTA TGTTTTTTTT ATCTGCTAAT ACTTCGTAAA TGTTTTGTTT TGAAACATCT 420

ACCCCTTGTT CTTCCAAGAG ATCTGAAAAA TAGCTAGTGG TTGATGCTTG TGGATCGGCG 480

TCAATTAGAA GAACTTTATA TTTTTTAGAC AAGAGTGTTG AAAAAATAAT AGCACTTGTG 540

CTTTTGCCAA CACCTCCTTT AATTGAGCAA ATGGCAACTA TTTTAGTGTT TTCTCTATCC 600

ATTTATTTAT AATTCCTCCA TCAGGCAATT CTTTGCAATA AAATTCATAC ACTTTTTTTT 660

CCAATCTGTT TAACATGTCA ATAAATGTTT GAAAATATTT TTTATTAATT TTTTCTTTTT 720

TGATTAACCT AGCAAGACTT CTTAAATAAC AAAACACACT GCCTTTTTTA AATTTAAATT 780

CTATATAATA TACTTTGGAA AATGTATATG ATTTTAAAGT GCCATTGATT TTGTATTTTA 840

TAACAATGTT TTTTATAGGT TTTCTGTATC CATAGAAAAT TCCTATAAAT TTATCATTTT 900

CTTTTGTAGA GAATAAATTA AGTCCTTCTA ATTTTCCTTG ATTAAATAAT TTTCTAAAAA 960

GAATTAAAAA TTTGTTTTTT TTATATCTGT TTATTTCAAA TTTGTATAGA TCCATTAGCA 1020

TTTTAGTGTG ATATATTGCT CTTTCGTTGA GAATTTCTTT TTTGATGAAA ATTTCTGGCT 1080

TTCTACTTTT TTTTATTATT TCTTTTTTTT TGTTTTTAAG TTTTTCTAGT ACACTTTTCA 1140

TTTCAAACTC TTAATTTATA TAGCTATTTT TATAAATATT TTGTGATTCT ATTAGTTTGA 1200

TAATTTCATT ATAGTATTGA TTATTAAATA TTTTTTTGTA TTCAAGTTTA TTTTGTTTGT 1260

TTAAGTATTC TTTAATTTTC GACCTTAAAT TTGTTGTGTT GGTTTTTCTA TGTAATTGAT 1320

CTAAAAGTAT GTTATATATA TTGGTTTCTA TATTGTCTTT ATGTTTTTGG GGTTCGGTTT 1380

TATTAGTATC GTCCTTTATC TTTTTTATAA TTTGATCGAA ATCCGGGGAT CCTCTAGAGT 1440

CGACCTGCAG GC 1452

(2) INFORMATION FOR SEQ ID NO: 69:

(i) SEQUENCE CHARACTERISTICS:
(A) LENGTH: 1426 base pairs
(B) TYPE: nucleic acid
(C) STRANDEDNESS: double
(D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 69:

CCCAGCCGGA AGGGCCGAGC GCAGAAGTGG TCCTGCAACT TTATCCGCCT CCATCCAGTC 60

TATTAATTGT TGCCGGGAAG CTAGAGTAAG TAGTTCGCCA GTTAATAGTT TGCGCAACGT 120

TGTTGCCATT GCTACAGGCA TCGTGGTGTC ACGCTCGTCG TTTGGTATGG CTTCATTCAG 180

CTCCGGTTCC CAACGATCAA GGCGAGTTAC ATGATCCCCC ATGTTGTGCA AAAAAGCGGT 240

TAGCTCTTCG GTCCTCCGAT CGTTGTCAGA AGTAAGTTGG CCGCAGTGTT ATCACTCATG 300

GTTATGGCAG CACTGCATAA TTCTCTTACT GTCATGCCAT CCGTAAGATG CTTTTCTGTG 360

ACTGGTGAGT ACTCAACCAA GTCATTCTGA GAATAGTGTA TGCGGCGACC GAGTTGCTCT 420

TGCCCGGCGT CAATACGGGA TAATACCGCG CCACATAGCA GAACTTTAAA AGTGCTCATC 480

ATTGGAAAAC GTTCTTCGGG GCGAAAACTC TCAAGGATCT TACCGCTGTT GAGATCCAGT 540

TCGATGTAAC CCACTCGTGC ACCCAACTGA TCTTCAGCAT CTTTTACTTT CACCAGCGTT 600

TCTGGGTGAG CAAAAACAGG AAGGCAAAAT GCCGCAAAAA AGGGAATAAG GGCGACACGG 660

AAATGTTGAA TACTCATACT CTTCCTTTTT CAATATTATT GAAGCATTTA TCAGGGTTAT 720

TGTCTCATGA GCGGATACAT ATTTGAATGT ATTTAGAAAA ATAAACAAAT AGGGGTTCCG 780

CGCACATTTC CCCGAAAAGT GCCACCTGAC GTCTAAGAAA CCATTATTAT CATGACATTA 840

ACCTATAAAA ATAGGCGTAT CACGAGGCCC TTTCGTCTCG CGCGTTTCGG TGATGACGGT 900

GAAAACCTCT GACACATGCA GCTCCCGGAG ACGGTCACAG CTTGTCTGTA AGCGGATGCC 960

GGGAGCAGAC AAGCCCGTCA GGGCGCGTCA GCGGGTGTTG GCGGGTGTCG GGGCTGGCTT 1020

AACTATGCGG CATCAGAGCA GATTGTACTG AGAGTGCACC ATATGCGGTG TGAAATACCG 1080

CACAGATGCG TAAGGAGAAA ATACCGCATC AGGCGCCATT CGCCATTCAG GctGCGCAAC 1140

TGTTGGGAAG GGCGATCGGT GCGGGCCTCT TCGCTATTAC GCCAGCTGGC GAAAGGGGGA 1200

TGTGCTGCAA GGCGATTAAG TTGGGTAACG CCAGGGTTTT CCCAGTCACG ACGTTGTAAA 1260

ACGACGGCCA GTGCCAAGCT TGCATGCCTG CAGGTCGACT CTAGAGGATC CCCaGAtGGG 1320

GTTATTATTG TTACTGTTAA TGACTATCTT GCAGAACGTG ATTCCAATTG GATGAAAGCC 1380

GGTTTTTGAA TCTTGTGGGG TGTTAGCGTn GGGGTTGTTC TAATCn 1426

(2) INFORMATION FOR SEQ ID NO: 70:

(i) SEQUENCE CHARACTERISTICS:
(A) LENGTH: 1425 base pairs
(B) TYPE: nucleic acid
(C) STRANDEDNESS: double
(D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 70:

TTTTGGTTGn AATTGCCACn ATAAAAGGGA TTCTTTTTTG GGTTTTATGG GCGGTATTCT 60

TCATTAtAAA nTTmnTCtAT ACAGCAAATA TGGAAAGACT TGAGAGTGCA TTAACCCCAG 120

CAATAAATGC GGCACTCGCT CCATTAAATG AAAAAATCAA TCAATGCATT GACTTAGTTA 180

ATTCTGATGA AAAAAATCTC AAAATATCTA ATGATCTGAA ATTCAATCAG GAAGGAAAAC 240

CTATCTATAA GGAAAGAAyA AATAATGCAA AATAACACTA TTGGTTTAGG ACTTAATTTA 300

CTATCCAGCT TAACTAACAT AGCTAAAACT GATACAAACA TAGATCATAA TTACATTAAT 360

ACTTTTAGTA AAGTAATAGA TTTTTTCTAC AAAACATATA TAAGCACACT AAAATCTATG 420

GAAACAGCTG AGTCAACTAA AATATTTGAA GAAATACAAG ACATTTTAAA ATACAACATT 480

GAGATAATAG AGGCTATCTC TACTGATAAA AGCAAAAGAA TTATCACTTC ACTTAAAGCA 540

ACACGTAACA AAATCATGAA AGAATATATC AAAATACTTA AAAGAGGTGA AAATGCTTAA 600

AAGATTGcAT TGTCTACTAA TTGCTTTGCT GCTATGTTGC ACCACTATTG CTAACCTACC 660

AGAAGAGCCA AAACCGCCAA TTATTCAAAC ACTAAAATCT TTAGCTAAAT ATGAAACACA 720

ACTTTCAGAG TATGTTATGT ACCTTGTAAC ATTTTTAGCT AAAACAAAAG TCAAAGTTAA 780

TGACCCAAAT TATCCAGAAT ATCCTTATCC AGACTTATCA ACACTAAAAG ACGAACACTC 840

CATAACTGCA GTAAAACACA ATATCAACAT ATATTTAGAG TACATTAAAA AAACAAAACC 900

AATAGCGGAA AAAGTCTATA ATAAATATTC CCAATTAAAA ATGTAAATTA CAAAAAGGTT 960

TTTCTTGCAA GAAATTCTAC TTTATAATTA AATTGGCTTT TACAACAGAA GAAAATCTAG 1020

ATATTAAATT TACTTTAATC TAATATCTAG ATTTTAACAT TTTCAACATG AATATTTACT 1080

AATTAATTAG TGCCcTCTTC GAGGAACTTT ATTACTTTGT CTATCTGTTC TACAGCGTTT 1140

TTAGACATTT TATCCCCATT ACCAGAAGTA TTGCTTcCAA GAAGTGGTAC AGTTACTCCA 1200

ACTAATTTTG CTTCTGACCA TATTTTTCTT TTTGAAATAT CCTGATCCTT GTCAGTAATG 1260

TTTTCAATGG TATTTTTAGC TGCTTTTAAC GCTGCTAATT TGGCCGCTAA TTTATCCTGA 1320

AAGGAATTTT GCAAATTTAA TAGTTTTTCT TTAAGCTCTG CATTGCCTAT ATCTTTTAAA 1380

TTTTCTTTTA ATTCATTTAT TTTGTCTTTA TAATCTTTAA ATTTG 1425

(2) INFORMATION FOR SEQ ID NO: 71:

(i) SEQUENCE CHARACTERISTICS:
(A) LENGTH: 1423 base pairs
(B) TYPE: nucleic acid
(C) STRANDEDNESS: double
(D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 71:

CAGGTCGACT CTAGAGGATC CCCTAAAATA ATTAACCCTG TCAATAAAGC GAGAGTGGAC 60

ATTGATTGGA TGATCTATTA CAGAAACCCC ATTAATATAC ACCTTTTTAA ATACAATCTC 120

TGTGTTATTT ATAACTTGTA TATTAAAACC TAAATCCATG TATCCGTCGT GATCATACCT 180

CAAAGTAAAA GATCCAAATT TGGTTTTTAA TTTAAGAATT AAACCGAAAT CATATCTATT 240

ACTATCATTT AAAATAACAA CTTCTTCAGA TATTTCACTA GGAAAAATAA ACCTGCTGGG 300

AATTAATAAA AAAGAAATTA ACAATATTAT TACTTTCATA CCTTTTATTA TTATACTATT 360

CTAAAGAAGA AATAAATAGA ATAAAAAATT TTAATTTCTC TTTTTTAAAA CTATTATTTC 420

TAGGTCAAAG ATTATAGAAT AGAAATAGTT TATTGCTTTA TCTAATGATA GCAGCTTATA 480

GGATCTTTTT TAAGATCGGT CTATCATCAA GAATATAAAT CACAAAAGCT TTTTAAAAGC 540

TATCTAAATT CTTTATTCAA GGCAATAATT TATGTAAAAA TAATTAAAAA AACCTCCATT 600

TTTGAGCAAA CATTTATACG AATTGATTTG GAAAGTCAAA TGCACAATTC TAACACTGAT 660

TGATAAAAAT ACTTTTTAAG TTTTTTATAT TCAAAATATA AAAAACTTAT TTATAAAAGA 720

TTTTTCAATA TCGATTTTTT TGTGATTTTA TTATTATTGG TATAAAATCA CATAGGGCCT 780

AACCATAAAT ACTCTTAAAG CAAGAATACT TATCTTAAGC CCTATAAATA GACATCGACC 840

AAAGTTAAGG ATGCTTATAG TTAATAGCAC CACTTACCAA GATTATACGC TATTATAGTG 900

TTAAAATCAA TACATTATTC TCAAATAATA TACATATTTA TTTATAAATT ATCTTTTAAA 960

AAATTTACTT CACTTTATTG ATTATTTTTC TAACACTTTC TGATTAAAGT CAATATTTTA 1020

CAAAGTATTT AAATTCGGGT ATTTGATAAA AATAGTGAAT TTAAATACTT TATTTTCCAA 1080

AAACTATAAT TTTATATTCT GCACACAAAA TTATCTATAT TAAATTTTTA ATTATATTTT 1140

TTACACTCCT TATATTCTTT CATAATTTCA TTAAGCAATT CTTCTTTATC TTTAAGTAAT 1200

TTTTCTAGCA AAAAACTAGT AAATCTTGCT TTTGATTTGT AATATATGTA TGCATCTTCT 1260

GTTTTAAGCT GAAATCTTAA TGGCCTGATA AAATTTCGAT TGGATTTTTT AACTTTCCCC 1320

CCTTCTTTAT CCTTTAAAAA AAATAAAGAA TTCTGTATAC CGTTTTCGAT TATGTATTTT 1380

TCCTGAACTA ACCCTTCTTC TATTGCATTT GCCATTCTTA AAT 1423

(2) INFORMATION FOR SEQ ID NO: 72:

(i) SEQUENCE CHARACTERISTICS:
(A) LENGTH: 1405 base pairs
(B) TYPE: nucleic acid
(C) STRANDEDNESS: double
(D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 72:

TATTTATnAA TATTGAGAAT ATTATCTAAT AAAATATTAA AGATGTAAAA ATTAGTTACA 60

AAAAATTGCT GTAGTGACAT AAATAATAAA ATTGAACTGC TAGAATTTTA TACAAAAAAT 120

AACAAAAACT TTATAAAGTT GATAATTATA AAAAATTTTA AGATTTTCTT GAAAAGTTTA 180

TCATATATAT AAAAAAAGAC AAGCACCATT ATTAATGTTT ATTAGTATAA AACCCCAAAA 240

TAATACAAAT TTAATCCCAA CAATATAGAT AGGATCTTAT TTTTTAGATA AAGTTTTTTA 300

AAACTTTAAA AAATATATTA AAATTTATAA AATATAAAAA GCCTATAATA CCGCACTTTT 360

ATTATCAAAA ATTGCTTATT TAATCTCATA AAAGCATCTT ATTGTTCTAT CAAGCTTATG 420

TATTCTCTAT TATAAGAGCA CAATTAATTA TACCAATTGG GGAGAATATT TTTATGAAAA 480

ACAAAATGAT TTTCTGTATC TGTGTTTTTT TACTTTTAAG CTGCTGTGCT GCAACCATGA 540

CACTGAAACA AAAATTGTTG ATAAAACAAA AACCTAATAT ATTAATGAGA TAAAAATTTA 600

ATAGCAGCAA GTAAAGAAAT CATCGAGAAA CGAACACTGC AACAAACTGA GCCAACAGAT 660

CAAGAACCTG TAGATAATAA AAACTGGGAG GAAGTTTTTG ATATAAATAA AAAAACTTAT 720

GACTTTATAA ATAGTTTTTT AACAAATGCT GAGTTCAATA TATTTGCAAC AATATTAAAT 780

AAACCAAAAC AATCACCAAG CAAGATGTTA AATAACATAG CAATTTTAGA GCTTAATCTG 840

GAAGAGACAA TTAATTACTT AGACTCAAAA AAAGATGTCT TAGATAAGGT AAACACCTTA 900

GATTTGGAAA AGATCAAAAA CTCTCTTGAA TAATTACTCT CTATAAGGAA TTTTTTTCAA 960

TAAGCATAAG AAAAACTTTA TTAGATCATC AAAATAATAC CGGTTCTATA AAAAAGGATT 1020

ATTCTAAATT AGATTCTTAT CTTAATACAA TACTTAATCA GTTTAATGAA AAAATTAAAG 1080

AGGTTGGAAA TTTGAAAAAA ATTATATTAT CAATAACTGT TTCAGCATTA TAAATTAAAA 1140

TTTATTAATG CAGGGGCTTA AAGTAAATTA AACCTTTAGA TATAAGGGGC TAATAAACTT 1200

TTTTATTAGC CCCTGTTAAC ATTCCTTTAA TCAAAAATAT TGAATTTTAA TTACAAAAAC 1260

AAAAAAACAA TTAGATTGTG AAAACAATAA AGATCTTATA TAAACAGATA TCAATGAGCT 1320

TAAATCCTAT GTAAGTAAAC TTGCCGATGA TTTAAACAAC TATCTGCAGA AGCAAGAAAT 1380

CTGCATTTAT AGTGTCAACA TATAG 1405

(2) INFORMATION FOR SEQ ID NO: 73:

(i) SEQUENCE CHARACTERISTICS:
(A) LENGTH: 1398 base pairs
(B) TYPE: nucleic acid
(C) STRANDEDNESS: double
(D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 73:

CnTCAACTTC AGCTCTGTAG TACTATCTGC ACTAAAACTA AGCTTGACTC CACGCTTGTA 60

TGGATACCCT TTAGCAGGAT AATTTTCTAT TTwATCTTTA CTACTAGTAA AAACTCCATC 120

CGAATTGGAG TAAATTAGAT TTTTATCTCT AAAATCTACA GAATTACTAA GCAAACCAGC 180

ATCTTGCTGA GGATTTTTCA TTAATGCTTT AATTTCTGCA ACTTTTTTAT CAAAATCTTC 240

TTTTATTTTT GTAATACCAT CGCTCATTAA AAACTCCTTT AAGCAATACT TGTTCTTTTA 300

TGTCTTTTTA GATTCTCATA AAATTGAGCT CGTCTTTGCT TGTAAGTATT ACTTATCGCT 360

TGTACAAACT CCGTGAAATT AATGGGCACA AAATTAGAAT CAAGCAAACT TGCTCTCTCT 420

TCTGATTTAA CAACAATATT GCCTCTGACA GAGTCAACAG AAGAAGAATT GCTACTCCCA 480

GTTTTTCTTA ATTTAATATT CACTTTTGCT AAAGAAACAA GTTGTTCTAG TATCTCTCCA 540

TCGATATGAC TTATATCTGA CGCTTTAGCA ATAGCTTTAA TTTGCTCAAT TGGAACAAAC 600

TTACGAACAA GCTCTCTACG TTGTGCCTGC ATAATGTCTT TTAGGGTGTA TCCTTTTGCA 660

AGTAACACTT CCTTGTTAAA ATGGTTGCTA AGATGCGCTT TTACAAGTGT ATCAATTTCA 720

TTAATTCGCT CAGCCTCTAG TAACAATTGC TTTTCAACAC GCTCTCGATC TTCAACTTCT 780

GCAAGTTCTT TTGTTATTCG CTCATTTATA CTCATCACGC CTTACCTCTT TAGGGGGTTT 840

ACATTGTCTG TTTGCTCTTT AAAGCGCATG TACTCTTCAA ATTCCTGCGC ACTTATAACT 900

TTAGTATCAG CCTTATTTTG CTGCTCTTCT TTATCTTGTG CTTGCAGGTC TTCTTTTTCT 960

TTTTTCTCAG TCATCTTTTA ACTCCTTTTC TCAAAATGAG AATAATTTCT CTTTTAAAAT 1020

CGCTAGCTCC TCATTATCAA AGGmGCTACT TTGTATAAGC TGGTTATATT TACTGTAAAG 1080

CTCAATTAGC TTTATATCTC TTTCCACTTT TTGCTCTTCA CTTAACATAA TCAGAGAATT 1140

AAACTcATAT CAAGCCCGAA ATACTTgTAA GTTTCAAGTT ACAAGCGTTC TCAACTTGTT 1200

CTTGcACACC CTTTAAAAAA TCGTAATAAT TACTCCTATC CCCTTTACCA TCATTTCCTA 1260

GCCCTTTAGC CTGTTCGTTA AAACTTCTGG TTAAGGGCTC TTtAGTATCT GCACCAATTT 1320

TtGCCTtAAT TAATGCTAAA GCCTCCTTTA AGTAACTAAG GTCGTATTTA ATAACCTCTA 1380

AACTAGCACT AGGGGTGG 1398

(2) INFORMATION FOR SEQ ID NO: 74:

(i) SEQUENCE CHARACTERISTICS:
(A) LENGTH: 1380 base pairs
(B) TYPE: nucleic acid
(C) STRANDEDNESS: double
(D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 74:

ATAATAGGCC CAATAAAGAA TATTTTGAAC ATGAATAATT ATTTAATTAC CAATATAAAG 60

CTAAATACAA AATAAATTCA ATATATCTTT AAATTCTAGA AAATTTTTAA TCTAAAAAAA 120

TCATTATTAT AGTGCCCAAA ACAAAATAAA ACTTAAACTG GGAAAATTAG TGTTACATAA 180

AATGAATAGG GCTTAAGACA AATTCTTTAT AAAAAACTTG CTTTAAGCCC TATTTCACGA 240

TCATATTGTG ATTCGAATCC GCGTCGAACT ATTTATAGTA TAACAAAAAT TAAATCATAG 300

TCAAGTTGTA TTTTAAATAA ATTTTAATCT TAATTACTAA AACTTTACAA TATAACTAAA 360

ATTGCTATAA GAGTATTTAC TTTTATAGCA ATTTTAGTTA TATTGTAAAG TGATCAATTG 420

TGAGGAGAAA CTTTTATGAA TTCAAAAACA ACAAATAAAA CCACTAGAAA TTGCTATAAT 480

AAAGTTCAAC ACAAATTAAT AGTTCTTATT TCAACAATAT GCTATCTAAA CAAAACACAT 540

AAGAAATATA CACAAAAAAC CATACTCTAT TATTTTAATA AAAATCTAAG AAAAAACGGT 600

CAACCTATTT CTACACTAAG AACTATGCAA AAGTATATTT ATAGACTACA AAAAGAAATA 660

AAAGTCACAA AAAACTACTA ACTATTTCTG ATAACAATAT CGATATCCAA ACCATGTAAT 720

AGAAATCCCA AACACATAGA GCCAGCCCCC CAAAATACAA GGAATTAAAA TCAAGCAACA 780

CATAAACCAT TTCAATAACT TCTAAGTATA AAATGCCAAA GCATAAATTT TGCTATTGCT 840

TAAACTATTT TCGAGTATAC ATCTGTATTA AATTTAAGCT TGTTTTTTCT ATCAATGAAA 900

TCATCTTTTG CATCCTTGTC AAATACAATT TCACTTTAAT TTTAATAAAA AATAAAATAT 960

TTGGACTCAC CAATAGGCTT CAGTGCCCGC ATTAAACCTA AATGTTTAAT TAAAAATTTT 1020

TGGATTGTTA TTCCCAATGC TTTTTCTATC TTGAAAGAAC TTTAAAAGTG CTTTCAAGAT 1080

AGTTTCTTTT TGCACGTTAC TTGAATTTAC ATTCAAATAA TAGGGCAAGT TGCTTTATAT 1140

ATACTCTTTA TTTTTTTTGA TTTTTGTATT TAAGTGTTGA TATTATTAGT AATATTTTAA 1200

ACTTTACTCT TAACTAAAAG CTTGTTTTAT TGTTAAAAAT AAAACACAAA CAATACCCTA 1260

TAAATAGTTT AATATTGCAA TATTATTTAA ACTATAAAAA TATGTAAATA ATAATTTATA 1320

AATTAATAAA AAACATATAA GGGAGCTTTC TTAATGAAAA TCAAAAATAT AGCAACATAT 1380

(2) INFORMATION FOR SEQ ID NO: 75:

(i) SEQUENCE CHARACTERISTICS:
(A) LENGTH: 1326 base pairs
(B) TYPE: nucleic acid
(C) STRANDEDNESS: double
(D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 75:

TnAGAGCAnG GATCCAGGAG AAGTTAAAGA TCCTATAATG TATAAATGGT TTTCGCCTGA 60

TCAAATTGAA GATGTTGATC TTCAGATGGG CTACCAAAGA ACTGTAAAGT GGGATGCGTT 120

TTTAAATGCT AATCCTACAA CGATTGCTAA TGAGGTTAAT ACTATTTCAA CTATTGGATT 180

TAGTTCTGAA GTGGTAAGAC TTAATTACTT AAAATTACAG TACAAATTTA GGCACTTAAA 240

GCAGGCGTCT GAAAAATTTT ACACTTCAGA CTCATATCTT GGTGACATTA ATAACAATTT 300

ACTTCCTTTC TCTCAAGCCT ATAAGCTTGC AAGTAGCGAG ATTATTAAAC TTATTAATCA 360

CTTTGTACTA ACAGGCACTG TTTCAATTCA AAAAGATGGA AAAAATCAAA AACGTCTACT 420

TCCCAATATG TATGGACTTC TTAATATGCC ACATCAAGTA AAAGAAGAAG TTGCTAGTGG 480

TGATAAAGAT AAAATGGATA AAATATTTGA AAAAATTGAG GCGGGACTTT CAAAGTTAGA 540

GCTAGGAGAC GAATTTTCTA CACCTATGAT GGTAATAGTT GACCCTACAA CTTCGCTCAA 600

ACTCGTAAAG CCATACGCAG CAGCACAGGG TGCAGCAAGT AGCTGCGAAA AATGGGAGGA 660

TGTTTTGATT CAAACTATCA AGGCTATTAA TAATAGAGAA GATGTCTACA TTGAAACTTC 720

AAACTTGCTA AAACATCAAA TACTTATTTA CCCATTAAAC CCAGAGCTTA TTAAGTTTAA 780

ACCTAGCAaG TATATGTTAC CTACACCAAA TGaACAAGTG GATAAAGATT CAACTGATAT 840

TGcTCATTCA TACATTGATT TTGTTTTAGG AGGGTTACTT GCTACTAGAA AAACTATTTT 900

GCAAGTACAT ATCAAACAAA GTTAAAAGTA TAAGGTAAGT GAAAATGAGT GAACAAGAAA 960

ACTTACAAAC ACAAGTTGAG GCTGAAGAAG AACTTTTGGT AACAAAACTT TATTCTGAAG 1020

TGTTATTGTT ACTAGGAATA GACAAACTTG CATTAAGCAG ACAAAATTTT CTACTTCATT 1080

TATCTTTACT TCAAGCTATT CTAGTAACAC GTGGTATTGA TGCTAGTTCA CTTACATATG 1140

AACAAATATT TTTACTTACC TTTTACCATA TGGGTTGTCA ATTAAGAAAA CAGGGAGTTG 1200

TTCGAGAATT TGAATTTGAT AGGATCAAAA AAGAGAAATT CAATGAACTT GAACTTGATT 1260

ATtATCCTAG TAGCAGTGGA GGCGAAGAAG GTGGCGAGGG GGGTTGTGGC TCAAACAAGA 1320

ATTTTT 1326

(2) INFORMATION FOR SEQ ID NO: 76:

(i) SEQUENCE CHARACTERISTICS:
(A) LENGTH: 1309 base pairs
(B) TYPE: nucleic acid
(C) STRANDEDNESS: double
(D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 76:

GGGCTGCTAG ATTAATTGCA GGTAGCAGTT GCTTGATCTG CAAAATTATC TATATTGCCG 60

CCGCTAAAAA ACCCTTGAAC GGTAGTTTTG AAGGTGTTTT TTCCTTCATC ATTTCCATTA 120

CATTTATCAA GTTCAGTCTT TATATGTTCA AGTGCTGATT TAATTTTGCC TTCATCATTT 180

TCTAAGAATT TATCAAATTT TCCAACACCA GTTAAAGCGG TTTTTAACCA GTCAAGTTGT 240

GTTTTTTGAT CATCAGATAG CTTTTCTCTA AGCAGGTCTT CTTTAGATTT AGATTTAGGT 300

TTTTCTTGTG TTGCTTCTTT TTGGGTTAAA TCACGCTTTT GTCTGCTTTT TGTCTGCTTG 360

GTATTTGTAT CATTAGAATT ACAGCCGTTT AGCATTAGTA AAAACAAACA AAATAATATG 420

TTGATAATTT TCATTGTTAC TCCTTTTTTT ATTATTAATA TTCACTTAAC TAAGTATTAA 480

TACTAAATAT GGGATAAACA ATTATTATTT GAATTGATAT GTTTTAAGTG AGGTAGTAGC 540

TATTTAGAAA TGAAAGCAAA TATTAGCCCG GCTATCATTG TGATAGACAT TGCTCCCATG 600

ATTCCTAATA CCCATTTAAG CATTTCTGAA AGAGACATTA AATTCTTTTC AACATTGTCT 660

ATTTTAGTGT TTAAATTCTT TTCTACAGTA TCTATTTTGG CATTTAAATT CTTTTCCACA 720

TTGTCAATCT TAGTATTAAG TTCGCTTTTA ATAGCATCAA TCTTAACATG TAAATTCTTC 780

TCTACGGCAT CAATCTTGAT GTCTAAATTA GATATAyCCT TTTGTAAATt CTTTTCTACA 840

GTATCTATCT TAGTATCTAA ACTATCTATT TTTAGATTTA AATTCTTTTC CACATTGTCA 900

ATCTTAGTAT TAAGTTCGCT TTTGACACTA TCTATTTTAG AAATAAGATT ATCAAATTTT 960

ATATCaAATT GTTTTTCTAA ATTTTCTAAA TCTCTATATG TTAGCTCATT GTGATAATAT 1020

CTTTTAGATA AATCTTGTGC TATTAGTTGT TCCATGCCCA GCCTAATAAA TTCTTTATAT 1080

ATTTGTTCTT GAGTTACACT TGCAATATTT GTTGACACTG TTTCCATAAA ATTTTCCCTT 1140

ATGGTCATAT TATACACTAT TTTAGATTGA TTGGCTTTAG AGATTTTTAT ATGTAAAGGA 1200

GAATTTCTTG CAAGAAAAAC CTTTTTGTAA TTTACATTTT TAACTGGGAA TATTTATTAT 1260

AGACTTTTTC CGCTATTGGT TTTGTTTTTT TAATGTACTC TAAATACTG 1309

(2) INFORMATION FOR SEQ ID NO: 77:

(i) SEQUENCE CHARACTERISTICS:
(A) LENGTH: 1300 base pairs
(B) TYPE: nucleic acid
(C) STRANDEDNESS: double
(D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 77:

TATCTATATC ACCATTTTTA AAGAATTCTG TAACCACAGT TTTGAAAGTG GTTTTTTGTT 60

GTTCTGCTTG ATCACCATTA CAACTATCAA GTTGAGTTTT TATATGATCA AGTGCTGATT 120

TTATTTTATC ATCATCATTT TCTAAGAATT TGTCAAATTC TCCAGCACCA GTTAAAGCGG 180

GTTTTAACCA GTCAAGATGT GTTTTTTGAT CGTCAGATAG CTTTTCTCTA AGTAGTTCTT 240

CTTTAGATTT TGGTTTTTCT TGTGTTGTTT CTTTTTGGGT TAAATCACGC TTTCCCCGTC 300

TTTTTGTTTG TTGGGCATTG TTTTTTAAAG TGTCATTATC ATTAGAATTA CAGCCGTTTA 360

GCATTAGTAA AAATAAACAA AATAATATGT TGATGATTTT CATTGTTACT CCTTTTTTTA 420

TTATTAATAT TCACTTAACT AAGTATTAAT ACTAAATATT GGATAAACAA TTATtATTtG 480

AATTGATATT CTTTAAGTGA GGTAGTAGCT ATTTAGAAAw rAAAGCAAAT ATTAGCCCGG 540

CTATCATTGT GATAGACATT GCCCCCATAA TTCCCAATAC CCATTTAAGC ATTTCTGAAA 600

GAGACATTAA ATTCTTTTCA ACATTGTCTA TTTTGGCATT TAAATTCTTT TCTACAGTAT 660

CTATTTTGGC ATTTAAATTC TTCTCTACAT TATCAATCTT AGTATCTAAA TTAGATATAT 720

CTTTTTGTAA ATTCTTCTCT ACATTATCAA TCTTAGTATC TAAATTAGAT ATATCTTTTT 780

GTAAATTCTT CTCTACATTA TCAATCTTAG TATCTAAATT AGATATATCT TTTTGTAAAT 840

TCTTTTCTAC ATTATCTATC TTGGTATTAA GTTCACTTTT AACAGCATCA ATCTTAACAT 900

TTAAATTCTT TTCTACAGTA TCTATTTTAG AAACAAGATT ATCAAATTTT ATATCAAATT 960

GTTTTTCTAA ATTTTCTAAA TCTCTATATG TTAGTTCATT GTGATAATAT CTTTTAgATA 1020

AATCTTGTGC TATTAATTGT TCCATGCCCA GTCTAATAAA TTCTTTATAT ATTTGTTCTT 1080

GAGTTACACT TGCAATATTT GTTGACACTG TTTCCATAAA ATTTTCCCTT ATGGTCATAT 1140

TATATACTAT TTTAGATTAA TTGGCTTTtAG AGATTTTTAT ATGTAAAGTA rAATTTCTTG 1200

CAAGAAAAAc CTTTTTGTAA TTTACATTTT TAACTTCAGA TATCAGTTTT AAATTTTTTA 1260

CTGTAgATTT TTTACAAAAA CAGTATTGCA AAAACTCTnA 1300

(2) INFORMATION FOR SEQ ID NO: 78:

(i) SEQUENCE CHARACTERISTICS:
(A) LENGTH: 1295 base pairs
(B) TYPE: nucleic acid
(C) STRANDEDNESS: double
(D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 78:

GAATTAATAA GCAGAGATGA TAATTTTTTA GGCGTTATTC ATGAACGTGA AGACTTGAAC 60

AAAAGGATAG CAGAAAACGA TACTTtCGAT TTAAATAAAG ATTaTATAAA AGAATATGaA 120

ATTACACTTG aAAAATTTTT TCAGTTGTCA AAAAAATTTT TAATTTCATA ATATATAGGG 180

AAATGAAATG AGTGTAAAAT TAAAACATAT GAATATAAAA ATAAAAGATc GTATTAATAC 240

TGGcAAAAAT CAAAAGCAAA TTGaAATTAA TTGTGATGAA GaTAAAATGG AACGATTTCT 300

ATTTTTAAAA GAAAGGCTAA TAATCAACTT CCAAAAAGAA ATTCACAATA AAATAGAAAC 360

AATGAAGATC TTAAAAGAGA TTAAAGATAA AGAATATTAT AAATTAGATG GCTATCAAAA 420

CTTTGAAATG TTTACTAGGA ATTACAAAAT AGCAAAAAGC CAGGCTTATG AATATTTAAG 480

AATrGCAAAT GCAATAGAAG AAGGryTAGy TyrGGArAAA krCATAATCG AAAACGGTAT 540

AyAGAATTCT TTATTTTTTT TAAAGGATAA AGAAGGGGrG AAgTTAAAAA ATCCAATCGA 600

AATTTTATCA GGCCATTAAG ATTTCAGCTT AAAACAGAAG ATGCATACAT ATATTACAAA 660

TCAAAAkCAA GATTTACTAG TTTTTTGCTA GAAAAATTAC TTAAAGATAA AGAAGAATTG 720

CTTAATGAAA TTATGAAAGA ATATAAGGAG TGTAAAAAAT ATAATTAAAA ATTTAGTATA 780

GATAATTTTG TGTTAAGAAT ATTAACCTAT GTGATTTATC AAATATATTA TTTAGATAAG 840

GTATTTGATA AAAAACAGGG AATTTCTTTG TTTATAAAGT AGTTATTTAA ATCATAAAAT 900

GTCAAAAAAA TCGGGGTAGT AAAGTAAAAA GTAAAAGATG ATTTACAAAT AAATATGCAT 960

ATTATTTGAG AATAATGTAT TGATTTTAAC ACCAACAAAA TTCATTTTCC AAATGATATA 1020

AAAGTTTTTA TAAATGAGCT TATAGGGTCG TTTTCAAAAT TAGGCTATTA TAAAGAGGCA 1080

AAAGAAACTT TGCAAAATAT TTTTTGTATA TTAGATAGTA ATTAAAAATG ATTTAGTCTA 1140

CTCAATAATT TATATGAGAT AATAAAGTAT ATTAAGGATA TGTGCTTTAT TAATAAAAAT 1200

AAAAATAAAA ATAAAAATAA AAATAATTAT ATCTTAATAA AAAAATTGGA AATACCGCAA 1260

TTATTGATAA TTTATCGAAT TTAGATACTA ATATA 1295

(2) INFORMATION FOR SEQ ID NO: 79:

(i) SEQUENCE CHARACTERISTICS:
(A) LENGTH: 1284 base pairs
(B) TYPE: nucleic acid
(C) STRANDEDNESS: double
(D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 79:

TATCnTTTTT CCTCAAAAGT TTAATTGGAA AATAAAAAGT TATnCTCTGG TGATATTAAT 60

AATTTTATTA TTGGGAATTC ACAACGCTCA GTTGAAGTTA ATGTTTTGGG ACAATTTGAA 120

AAGCTATGTA AACTTCTTAA AATTCCTTAT ATCCCAAGAC ATACAAATAA yTCATATATA 180

TTAATTGATT CACTtCGTAT TAATCTATAT GGAGGAGATA AGGCAAGTGA TTTTGAAAGA 240

TTTAGAGGCA GTAATTCGGC ACTTATTTTT GTTAATGAGG CTACTACTTT ACACAAGCAA 300

ACTTTAGAGG AGGTCTTAAA AAGACTTAGG TGCGGACAAG AAACTATTAT TTTTGATACT 360

AATCCTGATC ATCCAGAACA CTATTTTAAA ACCGATTATA TTGATAATAT AGCGACATTT 420

AAGACATATA ATTTTACAAC TTATGATAAT GTGCTACTTA GTAAAGGATT TATCGAAACA 480

CAAGAAAAAC TCTATAAAGA TATACCATCA TATAAaGCAA GAGTTTTGCT AGGTGAGTGG 540

ATAGCAAGCA CCGATTCAAT TTTTACACAA ATAAATATTA CTAATGATTA TGTATTTACT 600

AGCCCGATAG CATATTTAGA CCCAGCATTT AGTGTTGGmG GGGATAACAC TGCATTATGT 660

GTTATGGAGC GrGTTGATGA TAAGTATTAT GCTTTTGTAT TTCAAGACCA ACGACCAGCC 720

AATGACCCGT ATATTATGAA TATGGTTAAG ACCGTTTTAG AAAATTTTAA TGTACATACA 780

yTTTATTTAG AAGATAGAGA CAATACAAAA GGTGCTGGTG GATTGACyCG yGAATACATs 840

TTGCTAAGAA ATAATATGGG TCAATATTTT AGAATTGTTC CAGTTAAGCC AAAGTCTAAT 900

AAATTTAGCA GAATAACArC GTTAATTACG CCGTTTAyTT ATAAGAAACT kTACATTACr 960

AAGTACAGCA GTTCTTCTGT ATTTAATGAT ATTTATTCGT ATAAAGGAGA TAACAAAACC 1020

CATGATGATG CTCTTGATGC AATATCTGCA GCATATTTGA TGTTGTCTTT AGGGTATAGA 1080

GAGAGAAGTG TTCACTTTGG CAATCAAAGA TTTTTGTAAA TTTTATTGAC AAAAATAATA 1140

GTTTTTGCTA TCATACATCT AATTTAATAA AGAGAAATAA AAGGTGTGTG ATTTAAGAAA 1200

AACAAAATTA ATAGATAAGA TAAGTTCACT AGAACTATAT AAATACTCAA TATTTTTTAG 1260

GAATTATATA GAAAATGTAG CAGA 1284

(2) INFORMATION FOR SEQ ID NO: 80:

(i) SEQUENCE CHARACTERISTICS:
(A) LENGTH: 1271 base pairs
(B) TYPE: nucleic acid
(C) STRANDEDNESS: double
(D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 80:

GCTTGAAAAA TTTCTTTCGG GGCGCTTAAT GGAAGAATCA TTTCTTATTA GAATGTGGTT 60

AACTATTTTA AATTTTTTTC AGAAAGAATA AAAGCAAAGA AGAATATATA TAAAGCTTAT 120

GTTATAAAAA ACCTGGAAAA TCAAATTAAT GAAATGTTGT AAAAATCCAG TAATAGATTT 180

TAAGAAGAGA ATGTTTGCAT GTTGGTTTTG TGGAAATGTT TTTTAGTGTT TGCCGATATT 240

CAATGGAATT GAAAGAATTT TTTAGGAAAT TAGAAAAAGG TGGTATTGTT GTTGAGCAAA 300

CTATTTTAGA AATTATTCAA AGCAAAGTTC TTAACTCTAA GAACAATTTG GAAGAATTTT 360

TTAGATGAAG GTGAATATGA GCTTTTTTTA AAAAAAGAAA AAACCCAAAA CGAATTTAGA 420

AGAATCTCTT AAGGGTCAAG ATAAATGAAT ATATTAATTC TATTCCATCT AGTACTTACA 480

AAATCGTCTC GGATATGTTT GAGTTTTATT ATGTTTTTAA TAGTTTGGCG TTTTTCCCTT 540

ACAAATCTTT TTTTTCATTT TTTAATGTAG ACCTTTTAGA TAGTGCTGAG AATATTAGCA 600

TTGTTGACTT TGAAGGTTGG ATTTGGGGGG AATCCTCTAG AAGTCGACCT GCAGGCATGC 660

AAGCTTGGCA CTGGCCGTCG TTTTACAACG TCGTGACTGG GAAAACCCTG GGCGTTACCC 720

AACTTAATCG CCTTGCAGCA CATCCCCCTT TCGCCAGCTG GCGTAATAGC GAAGAGGCCC 780

GCACCGATCG CCCTTCCCAA CAGTTGCGCA GCCTGAATGG CGAATGGCGC CTGATGCGGT 840

ATTTTCTCCT TACGCATCTG TGCGGTATTT CACACCGCAT ATGGTGCACT CTCAGTACAA 900

TCTGCTCTGA TGCCGCATAG TTAAGCCAGC CCCGACACCC GCCAACACCC GCTGACGCGC 960

CCTGACGGGC TTGTCTGCTC CCGGCATCCG CTTACAGACA AGCTGTGACC GTCTCCGGGA 1020

GCTGCATGTG TCAGAGGTTT TCACCGTCAT CACCGAAACG CGCGAGACGA AAGGGCCTCG 1080

TGATACGCCT ATTTTTATAG GTTAATGTCA TGATAATAAT GGTTTCTTAG ACGTCAGGTG 1140

GCACTTTTCG GGGAAATGTG CGCGGAACCC CTATTTGTTT ATTTTTCTAA ATACATTCAA 1200

ATATGTATCC GCTCATGAGA CAATAACCCT GATAAATGCT TCAATAATAT TGAAAAAGGA 1260

AGAATATGAA T 1271

(2) INFORMATION FOR SEQ ID NO: 81:

(i) SEQUENCE CHARACTERISTICS:
(A) LENGTH: 1269 base pairs
(B) TYPE: nucleic acid
(C) STRANDEDNESS: double
(D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 81:

GAATTTTTCA AAATTACAAA ACTTTAACCC AAAATGGTAA AACTTTAATT TTTTGTAATT 60

TTTACATAAA AGTGTTAACT TTAAAATyCC AAACTTTATA ATTTTGGAAA ATTATCAATA 120

CTTTTTTAAT TTATTCTTTA TTTTCAAAAT AATCTTTATA TACTTATATA TTATGTATAA 180

GTCTGTAAAA GAACAACAAG AAAAAGGAAT AGATCATACA TGCAGAATAC TTATTCTTAC 240

CGAAACAATA TTTGAAATAA ATTTAATATT AGAAAATTAT TCTCAAAAAA CTCTACTCAA 300

AAAGTATAAC GAAAATCTCA AAAACAAAAA TCTACCTCCT AGTAATATAT CAACAATGAA 360

AAAATACTTA AATCAATTAG AAAAAGAAAT AAAAATCATA GCAAAATTCT ATTTTAAAAA 420

CGATCAATCT CTAATTTATT ATAAACTTAA TTATACCCTA GAAAAAATTT GGTTAAAACT 480

AATAGAATTA TTCTACAAAG AATTAAAACA ATTTATACAA AAGAACACTA CTACTTAATT 540

GTAAATACAT TATAAAATAA TCTTATTCAA AACTTTAGAA ATATATTGTT TTACGCTAAA 600

AAAATTTAAA AAATACTGTG CCATATTTGT AATATAAATT TAATATAATA GGGGGCTAAT 660

TCATTATGGA TGGAGTAaTT AACGATACAT TGGTCGCAAG AATGAAAAAG CAAATTAAAT 720

TTAATAAGAA TAAGTTAATC ATTCTTGTCA AAACACTAGA TCATATGAAT AAAGAATTAC 780

TTTATAGTGC AAATAAAACT TACAATTATG TCTTAATACA AAACAATTTT AATGAGGCTC 840

TAGCTAAAAC TTATCAACTT AGGGTTAATT ATAAAACCCT ATTAGAATAT CTTGAAATAT 900

TAGAAAAAAA TCCAAAAGTA ATCTTAAAAC GTCCCACAAA TAAAGAAAAT GAAAGCTTTA 960

TAGGCCTTTA TACACTCCTT TACCCTTTAG AAGATTGTTG CACTAAAATT TATAATTCTC 1020

ATCCTAATAT TTAAGCTAAT ATAATCCAGA AAATTATTTT TGCGAAATAG CATAATGCTT 1080

TAGAGTATTA AAGGCCTAAT AAAGAACAAT TAAAACTAAA AAATATATAA ATAAACCGTA 1140

AATTAGGAAA ATTAATGTTA CACCAAATGA ATAGGGCTTA AAACAATTTC CTTTAAAGAA 1200

ATTTCTCTAA GCCCTACTTC ATTGCTTATT ATTACGTCAA TTCGAGCATA AAGCCGAACT 1260

AATTTATAG 1269

(2) INFORMATION FOR SEQ ID NO: 82:

(i) SEQUENCE CHARACTERISTICS:
(A) LENGTH: 1225 base pairs
(B) TYPE: nucleic acid
(C) STRANDEDNESS: double
(D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 82:

CGGCCTTTTT TGTTGCAGCA TGATCAAGAA TCCCTTTAAA TTTAATGGTG AATTTGTCGC 60

TCACTAAGCC CTCACTTGCT TAAAATTAAT TCATACAATT CTTTTTCTAA TTTAATCTCA 120

GCAAGTCTAT TGACCTCTAA AgCTCGTCAT AAGGCAATTT CTTTACATTG TCGTACGAGC 180

AAaTATTCAT AATTACTGGA AAATAATATT TGTCGTTCTT AATCTCGTCA AGCAgTTAAA 240

ATATTTTTTT CTAGTCTCAT TAaGACTTGC AATAGCTTTA TCAATATCTC TATCTCTTTT 300

GCTCATTTAG CAACCAGCTC ATTGGAATTT GATGTACTTG ATGAAAGTGA AGTGGCTACT 360

TTTTCATAAT CAAAATTTTC ATTAATATAG TCAAAAGCAA CAAAATCACC AACATTATTT 420

TCATACTCAC TCAAATATAC TAAAGCGGGC TTTTTTAGAT CATTGTCTAA ATGAAAAGTA 480

TTAAATTGTG CAGTGTAAAT TATTGCAACA AGATAGTCCT TATAATAAGA AATAAATTCT 540

CTATTTTGAT CCAAAATCAC ATAGAATTCG TCTAAAAATT TTGGACTTAT CATTAAGCTT 600

GTGATTTCTC TTAAGTATTT AACCTCATTA AGCTTTAAAA CAGCGTCACT TTGATTAAAT 660

CCTAGCACCT TTATCCCATT CATAGACTGG TAATACTTTT AGTGGATATT CATAAGTTTT 720

ATTTTTAGTT AAAATTTTCA TTTTATATCT CATTATCATA ATAAGACTCT CCTTTTAAGT 780

GTTGTTTGGT TTAGTTTTTT GGCAATTAAT AGCCCTAATT TCAAAAGATA CTTTTTCGGC 840

CTCAGCAGAA TAACTTCTTG AAGGCTCTTC AGTAAAAATT GCATAGTTAG AAATAATTTT 900

GGTAGCAATT CTATCATTGA ATACTAAATC AAGCATTTTA TCCTCTTTTC TCACATCCAT 960

GTTGTAAAAC TGTTCATCAG AAAGTTCAGT TAACAAAATG TAGTCATGAC TACCTAGTGT 1020

CACTTCAATG TTGAAAACAT AAGTTATTGT TTTGGGATCT CTTAAGCTTA TTACAGGCAT 1080

ACCTTTATCT TCACTACTAA TCACTGCTCT TGTTGTAGGT TCGCTTGTAA GCTCTAGCTT 1140

GCCACTATGT AACTGCGGTA CCACCAATTG AAAAATAAAC TTCTCTTAAA TCATAAAATT 1200

GCATTTTTAG ACCCCCTTTT TAGCA 1225

(2) INFORMATION FOR SEQ ID NO: 83:

(i) SEQUENCE CHARACTERISTICS:
(A) LENGTH: 1200 base pairs
(B) TYPE: nucleic acid
(C) STRANDEDNESS: double
(D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 83:

CGGCTGGTGC TGTTAGTGCT GTTAGTGGGG AGCAGATATT AAGTGCGATT GTTAAGGCTG 60

CTGGTGCGGC TGCTGGTGAT CAGGAGGGmA AGAAGCCTGG GGATGCTAAA AATCCGATTG 120

CTGCTGCTAT TGGGAAGGGT GATGCGGAGA ATGGTGCGGA GTTTAATCAT GATGGGATGA 180

AGAAGGATGA TCAGATTGCT GCTGCTATTG CTTTGAGGGG GATGGCTAAG GATGGAAAGT 240

TTGCTGTGAA GAGTGGTGGT GGTGAGAAAG GGAAGGCTGA GGGGGCTATT AAGGGAGCTG 300

CTGAGTTGTT GGATAAGCTG GTAAAAGCTG TAAAGACAGC TGAGGGGGCT TCAAGTGGTA 360

CTGATGCAAT TGGAGAAGTT GTGGCTAATG CTGGTGCTGC AAAGGTTGCT GATAAGGCGA 420

GTGTGACGGG GATTGCTAAG GGGATAAAGG AGATTGTTGA AGCTGCTGGG GGGAGTGAAA 480

AGCTGAAAGT TGCTGCTGCT ACAGGGGAGA GTAATAAAGG GGCAGGGAAG TTGTTTGGGA 540

AGGCTGGTGC TGGTGCTAAT GCTGGGGACA GTGAGGCTGC TAGCAAGGCG GCTGGTGCTG 600

TTAGTGCTGT TAGTGGGGAG CAGATATTAA GTGCGATTGT TAAGGCTGCT GATGCGGCTG 660

ATCAGGAGGG AAAGAAGCCT GGGGATGCTA CAAATCCGAT TGCTGCTGCT ATTGGGAAGG 720

GTAATGAGGA GAATGGTGCG GAGTTTAAGG ATGAGATGAA GAAGGATGAT CAGATTGCTG 780

CTGCTATTGC TTTGAGGGGG ATGGCTAAGG ATGGAAAGTT TGCTGTGAAG GATGGTGGTG 840

AGAAAGGGAA GGCTGAGGGG GCTATTAAGG GAGCTGCTGA GTTGTTGGAT AAGCTGGTAA 900

AAGCTGTAAA GACAGCTGAG GGGGCTTCAA GTGGTACTGA TGCAATTGGA GAAGTTGTGG 960

ATAATGCTGC GAAGGCTGCT GATAAGGCGA GTGTGACGGG GATTGCTAAG GGGATAAAGG 1020

AGATTGTTGA AGCTGCTGGG GGGAGTGAAA AGCTGAAAGT TGCTGCTGCT ACAGGGGAGA 1080

ATAATAAAGA GGCAGGGAAG TTGTTTGGGA AGGCTGGTGC TGATGCTAAT GGGGACAGTG 1140

AGGCTGCTAG CAAGGCGGCT GGTGCTGTTA GTGCTGTTAG TGGGGAGCAG ATATTAAGTG 1200

(2) INFORMATION FOR SEQ ID NO: 84:

(i) SEQUENCE CHARACTERISTICS:
(A) LENGTH: 1182 base pairs
(B) TYPE: nucleic acid
(C) STRANDEDNESS: double
(D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 84:

GCTACAGAAA TAAAATTTTT AACAACAATT ATTTTTATTA TTTCGGCCTT TTTTCTCAAA 60

TTTTTATTGT CCCAAAAAGG TAACAATAAA ACGAATAACC TAGAAGAGGT GGCCAATTTG 120

TTATTTAAAA ACTTTAAAAT TTTTTGCAAT GCATTTTAAA TTCCAAATTA CTTTTGCATA 180

AATAAATAAA CTTCTCTTTG CGCAAATGGA AATCCAAACC GATCACTAAA ATATTTCTTA 240

AAAAATTATA TTCAGCTAGC CCAGATATCA TTTGCTTTTC ACACTGAAAA GCATCCTCTT 300

GAACACTAGA ATTAAAAACA CACTGTTTTT AAAATTTTCA ATTTAAATTA AAGATTGTAT 360

TTTGGCAAAA AATTTTCTAT ATCTATTTTA TATTCTCTGT CTTCATTAGA AATTAACATA 420

ATCTTTAATC GTTTTAATTT TTTGATTTTC TTAAAAAACC CTTTAGAAAT AGTAACACAC 480

AAGGTATTTT TTACTACAAA AGGAAATTCT AAATGAAAAA ACTTTTCATT TATATTCGGA 540

TCACCAATGT TAAAATCTTT TAATTGCTGC CATTTCTCAC TTGGTAAATT ATTTTCATGC 600

TTTGAAACTT TAGCATCTTC AAATCCTTCA AAAATTACAC TTTTAAAACC TAAAATTTTA 660

TTGTCATGCA CTTTAAAATC AAACCTATAA ACGGTAGATA ACGCTTTATA AGCATCTGCA 720

CGATAACCTG TCGCTTTTAT CATCTTTTTA TGCTTAAGCT CAGGAATTAT AAGCTGTGAT 780

TTTATAAAAA CCAATTCTTT CAAATTACTA TCTTTAAAAA GATATGTATA GGTTTCATCA 840

ACAAAAACAT TATCATCAAA CTTTTCAACA ATTTTATTAT AAATAACATT GTTTTCTTTT 900

TGTTTGTAAA AAAAGAAACC TAAAAAAATA CAAATTAATA GTAACCCAAA AAATAATATT 960

TTACCCATTA ATAATCTATC CTACTTATGA AAAATCATAT CAAATGCACT ATAAAAATGT 1020

GAGCTATTTT CCCTGCTTAC CCTGAAAGAA TATTTTTTTT CTCATTATTT ATCTCATCAA 1080

AACATTCAAC ATATACATCA ATTCCATTTT CTTCTGGCGA ACTTTCTTAT TTTATTCCAT 1140

TTTTTCACTG GAAATCCAAG GAAACTCATA TCCAAAATTT AA 1182

(2) INFORMATION FOR SEQ ID NO: 85:

(i) SEQUENCE CHARACTERISTICS:
(A) LENGTH: 1178 base pairs
(B) TYPE: nucleic acid
(C) STRANDEDNESS: double
(D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 85:

GnACnAGGAA CCGGCACTGA CGAGTCATTT AGACTGATTT TCAAGACCTT TTTAAATGTA 60

GACATTGAAG TTACTACTCC TGAAGCTGGG GTTATTGATA TCTCTTTAAA AGGGGTAATA 120

AAAACAAACT TTACTACATT TATTTCGCCT AGCACTAAGA AAGGAAAACG ACTAAAAAAG 180

ATAATTCTTA GAGAAAAGAA GCCGGGATAC GCTGCATCTA AAAAAGCTTT AGTATTTAAC 240

TCACTTCCTA AaGGCTAtGA TCATTCAATT TATGCTTTTA TTAAGAGAAT TATTCCTATT 300

GGTAGAGTTC TCAAAATTAA TAATACAGAT GGTAACAATA TTATTACTTT CAATAACTAA 360

GGAGGTTTTA TGGCTGATGA TCAAGAAAAA TTACTGATTG ATGAAGAAGA AACGGTTCAA 420

ATTAAAGATT TAAATAAGGT TACGACCGTT$AACAATACTG ATCTTTTACT GCTTGATGAT 480

GGAGCTGCAA GCAGCAATGC TATCACCTTT AAAAACTTCT TAAAAACCGT TAATCACCAA 540

ACATTTAAAG GCGAAGAGCT AGGCTATTTT AAAGAGATAA TTAAATCTAC AATCGCTACT 600

GAACTTGCAG CTGATAAAGA TTTTATAAAA AGCATTTACG ATTTAATCGT TGACAAGCTA 660

ATTGAGAATG AATCTAGTAA ACTTTCAAAT CTTTTTAGTA AAATCAAATC GCGCCTTACA 720

GATAGCATAT CATCAGCCAC TTTATCTAGA AGTGATGATC TTTTGATAAT GCCTTCATCA 780

GATACTATTC AAAAAACACC CGTTCCTAAA CATATACTTG GAGTACCATC AAATTTTACT 840

TATGGCAGCA TAACTAGAAG TACTACACTT TATCCTTCTG ACTATGAGAA TAAAGCGATA 900

TCTATTAATA TGGAAGACAA TGATGATGTA ACTCTTATTT TTTACAAAAA TTACGATAAT 960

GATCCCATTT ATCTGGATAT TGAGATTCAA GTAAAAATCA ATGGATAATA GGATGCAGAA 1020

AAAATCATTA AAACTTATGT nTTCTGATGA AATTACATAC AATTGGGTTT ATGAAATACG 1080

GGCCCTCGCG GACTATTCAC CAGAACTCCC ATTATAACGG AnGGTATATC CAAAAAAGAG 1140

CCTCCTGTAT GGAGATCGTC CCGATCTTTA AAACTGTA 1178

(2) INFORMATION FOR SEQ ID NO: 86:

(i) SEQUENCE CHARACTERISTICS:
(A) LENGTH: 1177 base pairs
(B) TYPE: nucleic acid
(C) STRANDEDNESS: double
(D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 86:

CATATGTATA ACAAAAATTA TTTTTGTCAG GCTTTTTACA GAAATTATTA TAATAAATAA 60

AAGCTTTATT AAATTCTCAT GTTAAAGAGC TTAAGAAAGC CGCTGGCTTA GCTCAATTTA 120

TTTTAGATAA ACTCACTAAA CAATTAAATT CAGTTCAATT TAGAATTAAA ATTATTTTGT 180

ATTTGTTAAA ATAAAAGAAC CTATTTAAAT TCTCTTGTTA AAAAATTCAA ATAAGTTCTA 240

CTTTAAAGCT ATATACTAAC TTATTACTTT ATAAAATTTT AATCATTCTT AATTTAAAAA 300

ATGCTTATTG AATATAGAAT AAATAATTGG AGCAAGCGTT ATTCCCATTA TTAAAATTAC 360

TTGTATTGTT CTATTACTTG CAGTAAGTTC GTTTTTTAAA ACATTTATTT TATTATCTAG 420

GCTAAATATA TCCyTTtGTA AGGTTTTTTC TACACTATCT ATTTTAGtAT TCAAGCTaGA 480

TATATCTTTT TGCAAAGTTT TTTCTACATT ATCTATCTTA GTATCTAAAC TATCTATTTT 540

AGAATTTAAA TTCTTCTCTA CACCATCTAT TTTGGCATTT AAATTCTTCT CTACAGTATC 600

AATCTTAACG TCTAAATTGG ATATATCTTT TTGTAAATTC TTTTCTACGC TATCAATCTT 660

AAAAATAAGA TTATCAAATT TTATATCAAA TTGTTTTTCT AAATTTTCTA AATCTCTATA 720

TGTTAGCTCA TTGTGATAAT ATCTTTTAGA TAAATCTTGT GCTATTAGTT GTTCCATACC 780

CAGTCTAATA AATTCTTTAT ATATTTGTTC TTGAGTTACA CTTGCAATAT TTGTTGACAC 840

TGTTTCCATA AAATTTTCCC TTATGGTCAT ATTATATACT ATTTTAGATT AATTGGCTTT 900

AGAGATTTTT ATATGTAAAA TAGAATTTCT TGCAAGAAAA ACCTTTTTGT AATTTACATT 960

TTTAACTGGG AATATTTATT ATAGACTTTT TCCGCTATtG GTTTTGTTTT TTTAATGTAC 1020

TCTAAATATA TGTTAATATT ATGTCTTACC GCAGTTATGG AGTGnTCGTC TTTTAGnGTT 1080

GATAAGTCTG GATAAGGATA TCnGGATAAT TGGATCATTA ACTTTAACTT TTGGTTTAGC 1140

CAAAAAnGnT ACCAGGnACA TAACATACTC TGAAAGT 1177

(2) INFORMATION FOR SEQ ID NO: 87:

(i) SEQUENCE CHARACTERISTICS:
(A) LENGTH: 1137 base pairs
(B) TYPE: nucleic acid
(C) STRANDEDNESS: double
(D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 87:

TCACGnCCAT GTtGtAAAAC TGTTCATCAG AAAGTTCAGT TAACAAAATG TAGTCATGAC 60

TACCTAATGT CACTTCAATG TTAAAAACAt AAgTTATCGT TTTGGGATCT CTTAAGCTTA 120

TTACAGGCAT ACCTTTATCT TCACTACTAA TTACTGCTCT TGTTGTAGGT TCGCTTGTAA 180

GCTCTAGTTT GCCACTATGT AGCTGTGTAC CACCAATTGA AAAATAAACT TCTCTTAAAT 240

CATAAAATTG CATTTTAGCC CCCCTTTTAA GCACTTAAGC TGTTTTGATA ATCAACTATA 300

TCTTGAGTAG TGATTACTAA AGCAACAGCA TTAATGCTAA AGTTATAAGT AATATTCACA 360

CTAAGTTCTA ATTTAAGTTG TGGTGTAGGA GAAAGGGTAA GGTTTAAGTT TTTATACTCT 420

ATAATCAATC CCCTGTCGAC AAACCTTTTA AGAAGACATT CAATTGCTGA AGTATATGCA 480

TTGTCTCTAG CACCACTAAG TTGTAGTGCA GATAATTTGC TATTTTGCCT ATTGTTTTTG 540

TTCCAAATTC TAATAAGCTC AATAATCGCT TCATTTTTTA TATAATGATA AGTGAAAAGC 600

TCGTCTATTG AACTTCCAGC AAGATCAACG CTCTCTTTAA AGGCAGGCAT ACCATCAAGA 660

CCAGTTTCAT TAAGAAGTGA ATAAAAGTTG ATTTTTGCAG TTCGCAACTT TCCAATTACA 720

GTATCATCAA CAAGTGGTGT AGCAGCCAGC GGCATGCCAT AAGGATTTAC AGCATGAAAA 780

ATACTAGCCT GATGTAAATA TTGACTTATA AATTTGAGGT GTAAATTGTC TTTATTATTA 840

CTGTAAACAG CAATATTTCT TTCTTTTTCA GTATTGCCTT TATCTTTAAA TAGTTCTTTT 900

ATTTCTTGTT CTTTAGTCGA GAATACAAAA AAAATTGAAG GTGTTTTAAA CTTATCATAA 960

TCATCTTTAT AAATCTTAAG TCCATCATCG GAATTATCAC CCTCAGTATT AATAAGTACA 1020

ACAAAAGTGT GTCTATGTAC TTTAAGATAT TTTTTTAACT CTTCGGGTTT ATCCTTATAA 1080

ATAAAAAGAA CAGCGGATTT TAATGATTCT TCACTTGAAT TGAAAAAATT TGACATT 1137

(2) INFORMATION FOR SEQ ID NO: 88:

(i) SEQUENCE CHARACTERISTICS:
(A) LENGTH: 1091 base pairs
(B) TYPE: nucleic acid
(C) STRANDEDNESS: double
(D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 88:

GTGATTTTTG AAAGATATGG ATGATTGTAG GTATTTATGA TGTCGCACTC AAAATTATTT 60

TCAGGTGCAT ACGCCTTAAA CCCTTTAAAT ATTTGAGTTA AATGATTTAA TACCATATCT 120

AAAGTGAAAA TCATTCAAGT GTTACCTTAT AAGTAATCTC TGATAACATT TTGGCTGTAT 180

CAACAAGTGG AATTGCTGCA GTGTTACTAC CCCTTTTAAA CTTACTTTTG ATTGTATTAG 240

CCTTTAAGGC TGGAGTGACT TGTGCTGATA GTAGATAATT TTCATAGTAC CTTATAAAAG 300

CTTGTCCAAT AGCCTCCATT CCCTATTTAG GGTCAAGATT AAACTTAGAA TTTATATAGC 360

TATTATTGAT ATATTCTCTA AACTCArAAC TACTAGcAAT TTTGGTTAAA TGTTTTCTTG 420

CTGGTAAATT GCTATTCCCT TTTTCATGCA TTTTAGCAAT ACCTGCACGA CCACCAAACC 480

ACCCAATTTC CAATTCCATT TTAAATTCTA GTTTGTCCAT ATAAATTCCT TTAAAACCAA 540

AGTAAAATAT CCGATTGAAG AGTCAATACT AAATATTTCA AAGTAAACTA AATCTGCAAT 600

TGATATTCGG TCTTTTAGTT CATAGTTAAG GTCTTGATAT GTGTAAAGTT TGGAATATCC 660

TTGAATATCA GACATATCAG AATCATAAAG AATTGCAAGT TCTTGTGGCT TTATGTCAAT 720

AATAACTCCT GCGAATTCAG TGTACTTATT TTTATCAAAT ACTCTCTGAT AAGAAGAATC 780

ATTTTCAAGC TTTACAACAG TACCTTTATA AAACTTTAAG GGTTTAGGAT CCTTAAATAC 840

GTTGATCATr CGAAATGACA TATCAGAAAG TCTTTTyCTA ACACCATTCA TTAGACAACy 900

CCCACACAAG ATGGCGTTGA AGTTTCTCTT TTTArTTTTT CTAAAAATGC ATCAAGTTGT 960

GAACAAAAAy TCTTGTTTGA GCCACAACCC CCCTCGCCAC CTTCTTCGCC TCCACTGCTA 1020

CTAGGATAAT AATCAAGTTC AAGTTCATTG AATTTCTCTT TTTTGATCCT ATCAAATTCA 1080

AATTCTCGAA C 1091

(2) INFORMATION FOR SEQ ID NO: 89:

(i) SEQUENCE CHARACTERISTICS:
(A) LENGTH: 1081 base pairs
(B) TYPE: nucleic acid
(C) STRANDEDNESS: double
(D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 89:

GGGAAATAAA TTCAAGAAGC AGGTATAATA ATTTTTATAA AAAAGAAGCA GATTTTTTAG 60

GTGCTGCTGT AGAACTTGAG GGGGCTTATA AAGCTATTAA GCAAACTTTA TTATAGATCA 120

CAAGGTATAA ATTTAAGGCT TAAGCCAATT TATCAAAAGA GAGGCATGTT TCTTGGTTTA 180

AAAGCATACT TTGCATACTT CTCTTTTGAG TACTACTATT TGAAAAGCTA TAAACTTTAA 240

CCTAATAGAA AAGCCAAATC TTTTTAAAAA TTTCTAAAAA ATAATTCTAC ATACTCTCCT 300

TATTACATTA AAAAATATTA TTGCTTATAT AAGGCACATA GTATAAAGAC ATTAATCAAA 360

ATTACCTTTT ACTAAGGTTT CAATCTCTCT AAACATGGAT AAGAACTACT TGATCAAGTT 420

ATAACAATCA AAACCCACTT TCTTTAAATA AAAAATCTTT TAATAATAAA ACCAAAATTC 480

CAGCCTTACT AAAGACCCTT ACTCTCTCGT GGATTTAATC TTCTTTATAT ATAAGGGTTA 540

GGCGTATCTA AAGATTTAAT CATTTTGAAT GATAGGGAAG AAGAATTCAT AAAGAATAGG 600

CAAAAGTGGT TTAGTTTACT GGAGCATATA CATTTAATTA TAAATAAGAA ACAATATTTT 660

CCATAAGAAC TGGAGTATAT AAATCATAAG AATAACTATT TTATAAAGAA TAAGTAAAAG 720

TAGTTTAGTT TACTGGAGTA TTTATCTATG TTAAAAGTAA ACGCACGGCG TATAAAGCCC 780

CTACTATAGT ATCCAATATT TTTTGAATTT AGGTCAATGT TGTTTAGTGT GTAAATAAGA 840

ATTATTAAGT GTGAAGATAG CCTATTTTTG CTATTCATAC TTAATATTTC TTTTAAAAAT 900

TTTTCCAAAT AGTGCCCCAG TAATTTTTAT TATTTATCAA ATAAAAATAT ATGTCTTTAC 960

ATTCATATTT CTAAATTGCT TAATTTGCAA AGAAATATTT TTTACGATTA AATAGTAGTA 1020

GGATAGTTTA GTTCTAACCG GAGTTTTAGT TTATCTGGTA TTGGTTGATA GTAGnCnTGT 1080

A 1081

(2) INFORMATION FOR SEQ ID NO: 90:

(i) SEQUENCE CHARACTERISTICS:
(A) LENGTH: 1078 base pairs
(B) TYPE: nucleic acid
(C) STRANDEDNESS: double
(D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 90:

GATCCATATG TGTCCCCTTT ATTTTTAAAT AAAAGATATA TATTTAAAGA CAGTTAGGCC 60

TCTTTTAGGC ATATTTTTGT TTAATAAAAA ATATTAAATT AGGGTTTATA ATTTTTATAG 120

ATGAAAATAA AATAGAAGAA TCTAATTTAA CTAAACAATT TTTGTTTAGT TAAAATGATA 180

TAGGGCTTTG CAAAGTAGAT ATAATTAAAG AAAATCTAAA ATCGCTAAAT AAAACTATTT 240

AAACTAAGCC CCATAATGAA AAAGTTTTAG TAAAAATATT AAAGAATATT TTTACTAAAA 300

TAAAAATTAA ACCAGCATTA ATAATACTTA CATTAGATGA TTAGCTACTT TTTTTAATTA 360

ATAAATTTTG CATTTAAAGT TCTATTCCAC TTATAAATAT TGACTATATC AATAATTTTT 420

CaAGCATTGG TACATTTTAT ATTCtAAATA TTTCGTTTTG TCGCTAATTT GTTGACATAG 480

GAATTATAAA AAGGCCATCA TCTTTTAAAT TAAAAAGTAA AATAATACTA ATAAATAAAG 540

ACCATCAAGC CCCGTCTTTT TTTTACTAAT AATACAATTG CATTGATTAT GGTTGTTATT 600

GATATTATTT TTTACTTTGA CAATGAATAT GAAAAAATTC TTTATTCTAA ATAAAGAAAT 660

TGGTATTGGT AATTGCAATT TATTATTTTA TTTATATTTT TTAAAAAATA TAAATAAAAT 720

ATAATAAAGA TTTATGGTAG AAAGCAAACA TCAAAAATAT TATTTTTATT CATTATTTTT 780

GTCAGAACTT GCAAGGACTT TGCCACATGC TGTATTAACT ATTATTTTAA TAAATAAAGG 840

GTTATCACTA AAAGATATTG CTATGGTACA AATTTGTTAT ATGGTAGCAA TTATTATTTT 900

TGAATTTCCA TCAGGTGTAA TATCAGATAT TTTTGATAGA AAAATTGTTT ACTTGGTGTC 960

AATTTTTCTA TTAaTGnCTT CTTATTTTAT TGTTGCTAAA ACCTCTTCAT TCGnGTTTAT 1020

TTGTGTTTCA GGTTTATAnA nGGGATGTCA GCnGCnATAG CACTGGCACG ATGACATA 1078

(2) INFORMATION FOR SEQ ID NO: 91:

(i) SEQUENCE CHARACTERISTICS:
(A) LENGTH: 1030 base pairs
(B) TYPE: nucleic acid
(C) STRANDEDNESS: double
(D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 91:

CCATTTTAAA AAATCAAATT TTACAATACA TTATTATTTG CCACCTTGTA AATATTTCAT 60

AAATAGGGCA TTCAAAATTG GCCCTAGAAT TGCTGCTGTT ATTATCATAA AATGTACTAG 120

TCTGTTACCC ATGCTAAGTT TTTGATTTAA ACTCTTTTCT ACATTGTCTA TTTTGATATT 180

CAAGCCATCC ATTTTTAGGT TTAAATTCTT TTCAACATTG TCTATTTTAG TGTTTAAATT 240

CTTTTCTACA GTATCTATTT TAGAGTCTAA ATTATCCATT TTTAGGTTTA AATTCTTTTC 300

AACATTGTCT ATTTTAGTGT TTAAATTCTT TTCTACAGTA TCTATTTTAG AGTCTAAATT 360

ATCTATTTTT AGATTTAAAT TCTTTTCAAC ATTGTCTATT TTAGTGTTTA AATTCTTyTC 420

TACAGTATCA ATCTTArTrT CTAAATTAGA TATATCCTTT TGTAAATTCT TTTCTACAGT 480

ATCTATCTTA GTATCTAAAC TATCTATTTT TAGATTTAAA TTCTTTTCCA CACTATCTAT 540

TTTGGCATTT AAATTCTTCT CTACACCATC TATTTTGGCA TTTAAATTCT TCTCTACACC 600

ATCTATTTTG GCATTTAAAT TCTTCTCTAC AGTATCAATC TTAACGTCTA AATTGGATAT 660

ATCTTTTTGT AAATtCTTTt CTACGCTATC AATCTTAAAA ATAAGATTAT CAAATTTTAT 720

ATCAAATTGT TTTTCTAAAT TTTCTAAATC TCTATATGTT AGCTCATTGT GATAATATCT 780

TTTAGATAAA TCTTGTGCTA TTAGTTGTTC CATACCCAGT CTAATAAATT CTTTATATAT 840

TTGTTCTTGA GTTACACTTG CAATATTTGT TGACACTGTT TCCATAAAAT TTTCCCTTAT 900

GGTCATATTA TATACTATTT TAGATTAATT GGCTTTtAGAG ATTTTTATAT GTAAAATAGA 960

aTTTCTTGCA AGAAAAACCT TTTTGTAATT TACATTTTTA ACTGGGAATA TTTATTATAG 1020

ACTTTTTTCG 1030

(2) INFORMATION FOR SEQ ID NO: 92:

(i) SEQUENCE CHARACTERISTICS:
(A) LENGTH: 1028 base pairs
(B) TYPE: nucleic acid
(C) STRANDEDNESS: double
(D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 92:

TTTTGTTGAA TTTCCTGTTA AATTTTCTGA ATTGGTGTGA TTGCTTGTGT TTTTTAGATT 60

TCTAGAATTG TTGCTTCGTT TTGTTTTTTT TAGACTTTTA GAAGTGGTAG GATTTTTTGG 120

TTCGTTTGGG TTAACATTGC CAAAAGGTGC ACATGATATG CAAATTGAAG TTAATATTGC 180

TGTAATAACG TTAAGTTTAA TAATATTTAA TTTAAAGTTT TTCAAAATAT TCTCCTTATA 240

AATTTGAATT AATATTTATT AATTTTAGTT CAAATATATA ATATTACAAT TTAATATCAA 300

TATCAAATAA GTTTAATATT ATTTGATATT GAAAATTTAA TTTCTATTGA TGTTTTTAGC 360

GTGGATTTAG ATTGCATGAA TTTTAAAAAT AAAAGTTAAT TCTTCTCTTT TTAAAATATG 420

AAGTGTAACA ATTTGTTTGG ATTTAATGGG TTTAATCTAA GGATCAAGAT GAGGAATTTA 480

GAAATTATAA CGAACTAAAA GAACAATTAA AATTAAATTT GAAATCTGAT ATTAATAATA 540

AAATTCAAGA AATGAAAATT CTACACGAAA TTAAGCAAAA ATAACTTTAT AAATATGACT 600

GTTTTAAAAG TTTTAAGCAG TTTATAAAGT CTTATGTAAT TGCCAGAAGT CAAGTGTATA 660

TGTATTTGAA AATTTATGAG AAAGTTTTAG AAGGGTTTAT TTCTATTGAA AAAGTTAAGG 720

AAATGGGGTT TGTAGCTGCA TATAAAAATA TACTAAAGAA CAACTCGTCA TATGTATATA 780

AAGAAAACAT GATTGAAGAA AATATAGCAG AAGATGGTGA TAGTCAAAAT ATGTCTATTA 840

AAATTTTAAT TAAAGATAAA GAAGTTTATG ATTTTTGCAA AAAAGATACT AAAAGAATAT 900

CTTTTATTTT AGGGGGGTTC ATTAAAGCAT TATTGAATTA AGTTGGAGAA TTTTTCTTTT 960

TGTATTTTTA TTAGCAATAT ATTTTCCCAT AGAGGCTTTT TGTGTCTACT AGAATAGGTA 1020

ATAAGATT 1028

(2) INFORMATION FOR SEQ ID NO: 93:

(i) SEQUENCE CHARACTERISTICS:
(A) LENGTH: 1002 base pairs
(B) TYPE: nucleic acid
(C) STRANDEDNESS: double
(D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 93:

AATTTTTnAA AAAAAAAAAT TTTTTTTTTA AACCCCCAGG GnAAAATTTG GGGAAAACCC 60

CCCCAAGGCC GTCCTTTATG GGTCCTGCCT TTAGCCCCAG AGGACGTTAA AATTGGATTT 120

ATTCAAACTT TTTATAATTA AAATTGGCAC TGTAAGGAGT TTAAACAACA AACTCAGAGG 180

CTATTGTTAC AATACCCGAA TTTGAAGATT TAGAAATTCA CACAAAAAAT ATCTCTAATA 240

TCAGTTTAGA ATTATCAAAA GGTGATAACG TATTGCTACT TCAATCAAGC GTTAATATTT 300

TTGATAAAAA TAACGATAyC CACTTTGACA AACATCATTT TTATATACTT AGTGCAATTA 360

GCCCAAAGAC TTTAAATCTA ATTTCTGATA CTGTTAAAAT TAAAGCAAAC AATAAAATTG 420

AAATAGCCAA TGAAATAACT TCCTTAAAAT CAATTCTAGA GAGTATTGTA AGTGCTATTA 480

ATGGAATTAC TGTAAAAGGA CAAGCGGTCG TTGACTATGC AAGCTTACAA ATAGCAACAT 540

CTAGAATTAG CAATAATATT AATAGTTTGT TTAAGTAATT TTTGCTAATT ATGGTATAAT 600

TACTAGTATG GATTTAAGAT TAGGCAATAA TTTTGAATTG GTATTTAATA AAGATATATC 660

ACTTGTTGAT GGAATTGATG AACAAAAACA AAGaTTTTTG ATATTTTTAA AAACCTTAAG 720

GGGTAGTTTA AGCTATGCTC CTCATTGGGG ATTGGACTAT TTCTTACTTT TAAAACTGTT 780

AAAAATTAAC AATCTTCACG CTGTAAAAAA TTATTTTCAT GAAATATCTA AAGAGCTTAA 840

CTTAGrTTTA ATAAATATTT CAACTACTAT ACAAGACAAC AAAGCACATA TATCCyTTTT 900

TTTCTCGGGC GATGTTTTAA ATATGGAGtT TAATTTaTGA GCtAGTTTTT GATTCTGrTT 960

TgGcwTTTAA AACGTACAtT AAGGgTATTG TAAGAGCTAA AA 1002

(2) INFORMATION FOR SEQ ID NO: 94:

(i) SEQUENCE CHARACTERISTICS:
(A) LENGTH: 998 base pairs
(B) TYPE: nucleic acid
(C) STRANDEDNESS: double
(D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 94:

GGGCATTATG TACTGATAAT GATGATGCTT TAGAAGATCT TTTTAAAAAG AATGCTGAGC 60

TTAAGAGTAT AGAATATTGG GTAAATATTT TAAAAAAATA TTTCAATAAA ACTAATAGAT 120

TTGATGATCT AAATAAGCTT AAAGTATTTA TGTCTGATAA TCGAGACGTT TATAAAACAA 180

AAGTATTAAA ATTCTTTTGC ATGTTGAAAA AAGAAAGACA ATTTAATTAT ATATTTGCAG 240

CATAGCAATA TTAAAGCCCC CAAATAGGGG GCTGTTAGCT ATTAGGAACC ACCATTGTTG 300

CAGTTACTAA CCGCATTGTT TGCAAAATTA TCTATATTGC CGCCGCTAAA GAAACCCTGA 360

ACTGTTTGTT TGAAGGTGCT TTTTTGTTGT TCAGAATTAT CCCCAGTACA CTTATCAAGT 420

TCACTCTTTA TATGATTAAG TGCAGATTTT ATTTTGCTTT CATCATATCC TAAAAATTTA 480

TCAAATTCTC CATCATTACC CAGAGCTTCT TTTAACCAGT CAAGGTGTGT TTTCTGGTCT 540

TCAGATAGCT TTTCTCTAAG TAGTTCTTCT TTAGATTTAG GTTTTTCTGG TGTTGCTTCT 600

TTTTGGGTTA AATCACGCTT TCCCCTGCTT TTTGTTTGTT GTGCATTGTT TTTTAAAGTG 660

TCATTATCAT TAGAATTGCA GCTATTTAGT AGTAGTAAAA ATAAACAAAA TAATATGTTG 720

ATGATTTTCA TTGTTATTTC CTTTCCTTAT CTCCAGTACA ATATGTTGAG TAAAAATAAA 780

ATTTATTCTT GTAATTATAG AGCTTATTTT TAAAAATCTT TAAAAATATT AATTGAGAGA 840

TTTATATTTT TCGAATGTTG TGCTAGCnTT TATTTCATTA TTATTGAATA TAGGAGTAAC 900

TAATGAGAAA TAAAAACATA TTTAAATTAT TTTTGCATCA AGGGATTTGT AATGGCTGTA 960

AAGCATATGT AGAAGAAAAG AAAGAAATGA TCATAATG 998

(2) INFORMATION FOR SEQ ID NO: 95:

(i) SEQUENCE CHARACTERISTICS:
(A) LENGTH: 996 base pairs
(B) TYPE: nucleic acid
(C) STRANDEDNESS: double
(D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 95:

AAAAATATTT TTATTTATAT TTTATTGTAA TTATTCTTAA ATGATATATA ATATCAATTA 60

AGAATAATTA TTATTTATAA TATATATTCC TACTTAGATA AAAGGAGATA TTTTTATGAG 120

AAAAAGTTTG TTTTTATATA CATTATTAAT GGGAGGATTG ATGTCTTGCA ATTTAGATTC 180

GAAATTATCT AGTAACAAAG AACAAAAAAA TAACAATAAT GTAAAAGAAG TTTCGAATAG 240

TGTTCAAGAA GATGGTCTTA ATGATTTATA TAGTAATCAA GAAAAGCAAA AAAGCTTTAC 300

TAAAAATTTT GGAGAATGGA AATATGAGGA TTTAATTAAT CCTATAGAGC CTATAATACC 360

TTCAGAATCA CCAAAGAATA AGGCTAATAT ACCAAATATT TCAATTGTGC ATACTCAAAA 420

AAAAGAGATA AAAGAGGAGG ATTTAATCCC TTCTACTAAT GAAGAAAAGG AAGCTGATGA 480

AGCAATTAAA TATTTAGAAG AAAATATTCT TCAAAACTCT AAATTTTCTG AATTAATTAG 540

AGAAGTACGT GTACTTAAAG ATGAATATGC TTTAATAAAC TCTGATTTTT ATGATGTAAT 600

TGAAAAGATT CACAATAAAA AAACATCATT AATGGAAAAT TATAAGAACA ATAGAGATAA 660

GATAAATAAA TTAACACTGT TGCAAAATAA TTTAAAGATA AATATTGAAC TTGAGCAGCT 720

TATAAATATG ATTGATATTG CAGAAAATGA AATAAGATCT GCGGCTTTCT TTTTTGACAC 780

CGCTCAGAAA AGGTTAAAAG AAAGTATTAT TAAAAGATTA GAGAGTAAAA ATAATAGATC 840

TTATTATGCA TTAGAATTGT CTAGACAGGC TTTAAGTGAC GCAAGAAGTG CTTTAAGCAG 900

TTTAGAATCT TTTGCTTTTA AAAGAGCTGA ACCAATGGTA AGAAAGAAAA AAATAAAAGA 960

GCTTATTAAA CATGCAAAAA CTGTTTTAGA AAGTCG 996

(2) INFORMATION FOR SEQ ID NO: 96:

(i) SEQUENCE CHARACTERISTICS:
(A) LENGTH: 986 base pairs

(B) TYPE: nucleic acid
(C) STRANDEDNESS: double
(D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 96:

CGGACTATAT ATACTAAAAG GGACTTGTGC TTGTATTCTA TTGGCTAGTT GTTCTTTTAC 60

AAGATGAGCG TCAAAGCGTA CATaGTATTG CCAAAAACAG TCATTTTGAG TAAAGAAATC 120

ATATCCTTAT ATAAGGATGT TAGTATTCCC CCGTTTAATG AkATATGTTC ACCAGTCATT 180

ACCGGATTGT AGCTTACATA TTCCGCTTTT CTATCATAAT AATTGATAAC TGGTCTTTTA 240

GAACAACTAG TATTATAAGT GCGTGTTATG AGTTCATTTT TTGGTTTTAT AAAAAACAAT 300

TGAGGAATAT ATCCAAAACC TTTTAGATCC ATTCTAGGAA ATAACACTAA AAAATTATCT 360

GcTCCGAAAA GGGCAAATAT TTGGGTTATt ACATCTCTTA TTATTCGAGT AATCTCCCTG 420

ATTTCTTcTT TtCAATATCA TTAATTTTTT CCTTGATTTT TTTCTTTtCA ATATCATGAT 480

TGTTAGTAAT TTTATTATTA ATATCTATTT TGTTAGCTGC ATTGTTAGCA ATTTTTTTGT 540

TACTTGTCAT AAGTAATTAC CTTTTGTAAA AATTATGGTG TGCTGTTAGC ATTGTCTgAT 600

TTTGAATTTC TTCTTGTAGT TTTTTTAGAG CCGCACCCTC ATCTCCGCCC ATCCATCCAG 660

GTAGCATCGA TTTtAATTTk GCAAAGAAAT AATtAAGATt AAAAATACTT TtAATGCCAT 720

tAATTtAtGGG ATAATAAtGT GTGTTTCAAA CGCAAaGTCT TtAAaGTAAT aGTTATCTTA 780

TAAGAGGTCA AGTAAGGGTC CAAGACAGTA GTGGTTAAGT TTTGAAGAGT TTGCTCAGCT 840

GAGGCCAAAT TACTTGAATA CTCTCAGCAT ATTGACTTTT TTGTAAGGCC GAAAGATTAA 900

AATCCTCGAA CATTTCCATC ATCTGGTAAn TCTAGACTCT AGAnCTACTT GTGCCCCGCT 960

TTGCCAGGCC ATTTGGCATC TTCTAT 986

(2) INFORMATION FOR SEQ ID NO: 97:

(i) SEQUENCE CHARACTERISTICS:
(A) LENGTH: 976 base pairs
(B) TYPE: nucleic acid
(C) STRANDEDNESS: double
(D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 97:

ATTGAGCACT CCTTTACATA TTCATCAAGC TCGCTTTTTA AAGAATTAAT TTCTCCATTA 60

ACAACTTGCT TGTTTTTTTT ACTACTTGCT TTATTTAAAG CGTCAATTTC GGCTCTTAAA 120

TTTTCTATTT TAGTATGCAT ACTAACAAGC TCAACACTAG AATATTGCTT AAATGCATTT 180

ATAAATCCTA ATTCTAAATT AGCCCGCTCT AAATCCAATT CGCTTCTAAC TTTCCTAGCG 240

TTAACTTCTG ATCTAAAGGT TTGCGACAAA AGGTGTTCAA AAGTATCTTC ACTAATTGTT 300

ACTCTAGAAT CCTCGCTAAC AGAAGTTTCT CCACTTTCCC ATTTTTGTCT CATTCTCCAC 360

ACATTTACCC TAGAAACTCC CAATTTAACC GATATTTCTC TATCATCTAA CGATCCTTCT 420

CTAAAGTATG CAACATAATC ATtAAAAGAC CTTTTAGCTC TTTTCAAAAC AATTTCTCCT 480

AAAATAACTA AATTAACAAA TTGTTACTCT AAATAGTAAA TCAATTTGTT AATTGTTAAC 540

ATTAACTATT ATCTTATTGA TATCTATTGA CAGGTGTTTG GTATTTTTTT GACTTTTATT 600

GATTTAGAAA TAGCAATTAA CTAATTTATT GAATTTTGCA ACAACTTGAC TATATAAATT 660

AGGGAAAATC TTTTATTGTT TTAATTAGAT CATCGCTTGT AAAAATTCTC TTATCATAGT 720

TGTGkATCCT TATAAATAGT ATATCTTTAA ATTCGTTGAT CATAATTAAT TGATATTGTT 780

TTGAAACTTT TTGATAAATA TGGTTAAGAA TACCATAAAC AGCCCCCAAA AATATCATGG 840

AATCATACTC TCCTAATTTT TTCAAACATT TCTTTAGCAT CCCTTTCTTG TCGCTATAAT 900

CAACTTGCAT ATTTTTGGAA TTTTTATATT TTTnTATTAA ATATTTATTT TTCAGAACGT 960

CTTTAATAAT TTTnTT 976

(2) INFORMATION FOR SEQ ID NO: 98:

(i) SEQUENCE CHARACTERISTICS:
(A) LENGTH: 968 base pairs
(B) TYPE: nucleic acid
(C) STRANDEDNESS: double
(D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 98:

GTCAGGCTTT TTACAGAAAT TATTATAATA AATAAAAGCT TTATTCAATT CTCATGTTAA 60

AGAGCTTAAT AAAGCCGCGG GCTTAGCTCA ATTTATTTTA GATAAACTCA CTAAACGATT 120

AAATTCAGTT CAATTTAGAA TTAAAATTAT TTTGTATTTG TTAAAATAAA AGAACCTATT 180

TAAATTCTCT TGTTAAAAAA TTCAAATAAT TTCTACTTTA AAGCTATATA CTAACTTATT 240

ACTTTATAAA ATTTTAATCA TTCTTAATTT AAAAAATACT TATTGAATAT AGAATAAATA 300

ATTGGAGCAA GTGTTATTCC CATTATTAAA ATTACTTGTA TTGTTCTATT ACTTGCAGTA 360

AGTTCGTTTT TTAAAACATT TATTTTATTA TCTAGGTTAA ATATATCCTT TTGTAAGGTT 420

TTTTCTACAC TATCTATTTT AGTATTCAAG CTAGATATAT CTTTTTGCAA AGTTTTTTCT 480

ACATTATCTA TCTTAGTATC TAAACTATCT ATTTTGGCAT TTAAGCTCTT TTCTACATTG 540

TCTATTTTGG CGTCTAAACT ATCTATTTTA GAATTAAGTT CATTTTTAAC ACTATCTATT 600

TTAATATTTA AATTCTTCTC TACATTATCT ATCTTAGTAT CTAAACTATC TATTTTGGCA 660

TTTAAGCTCT TTTCTACATT GTCTATTTTG ATATTCAAAC CATCTATTTT TAAATTTAAA 720

TTCTTTTCCA CATTGTCTAT TTTGGCATCT AAATTAGATA TGTCTTTTTG CAAATTCTTC 780

TCTATATCAA TTATTTTCTC TTTTAAAAAT TCAAAGTTGT AATaTCATTA TGCAGAAAAA 840

CAAAATCTAT gCTtCCTgCT AAACCCTATA TTAAAAATTC GTTTTTAATA CCTTTCTAAT 900

GGTTAATAAT GGTTTGGTAA TGGCCTAAAA TTGGTTCCAT AAGGATTAAC CCTTTTAAAT 960

GGTTTATA 968

(2) INFORMATION FOR SEQ ID NO: 99:

(i) SEQUENCE CHARACTERISTICS:
(A) LENGTH: 954 base pairs
(B) TYPE: nucleic acid
(C) STRANDEDNESS: double
(D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 99:

GCAGGTCGAC TCTAGAGGAT CCCCTTTAAC TAGAATTTTT CAAAATGATA AAACTTTAAC 60

CCGAAATGAT AAAACTTTAA TTTTTGCAAT TTTATTCTCT TGTTTTTTTT AAAACGATTA 120

GAATAATCGT TGAkCAGGTT TATTGATTAT CAATAAACCT GATCTATAAT ATTATAAGCG 180

GTTTTTGCAA GTTTAATAGG AGCTATAATA TCCATGAACA AATTATTGAT ATTCATTATT 240

TTATTAGTCT TTTCATGTAA TTTAAGTAAT TCTGATCAAA ATAATCCACT AAACATGTCA 300

AATAAAGAAA AAATAAGCGA ATATCAAATA AATGAGTCGT CAAACAAATA TTCAATTTTC 360

AAACGAAATT CAAGCGTTAA AAGATACACG TTCAACCATT ATTACTAACC AAAATGATAA 420

TATTAATTCT ACTATTAACT ACCCACCTTA TATTCAAACT ATCTTAAAAA TAGAAAAACA 480

AGTTGACGGA AATATTAATG GGATGACTAA AGAAAGTGGC ACAGAAACTA AAAAGCTTTT 540

AGAAATTCTA AATGGGAATA TTTCTCGATT TAAAGATGCA ATTCAATATG GAGGAAGTTT 600

TAGGGCTAAA GATGTTAGAG AAAATCAAAC CCAAAAAGAA AACAACAAAG ACTCGCATAT 660

TCATGTCGAC GATTTTAAAG AATACATACA TTTAATCATG CCTAGCATTA cAATAATGCT 720

GATAGTAGTA GTAGTTATTA CTATACCAAC TACATAATAA ATGGAGACAA TTTGTkAAGA 780

ATTATTAGCA ACTTATAArA AATCTTTATA AATTACCAAT ATTCTTGACA ATTTTAATAC 840

TATTTTTTTT ATATACTATA ATATTATGAA AAAAAATCAA AAAAACAAGT GCTCAGAAAT 900

AGAAAAAACA CAATTAGAAA TAATAAATAA CCAATCAGAA ATAGAAAAAC AACG 954

(2) INFORMATION FOR SEQ ID NO: 100:

(i) SEQUENCE CHARACTERISTICS:
(A) LENGTH: 946 base pairs
(B) TYPE: nucleic acid
(C) STRANDEDNESS: double
(D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 100:

AGAAGAAAAT AAATTATACA GATCTTCTGT ATCTTTTAGA TATTTTTTTT ATGATGAAAA 60

TACAAAAAAG AAATTAGGGT ATAAAAAAAT AATAACAATT TTCAATTTGC TTGATAAAGG 120

AAGTGATGCA ATAAAGTTTC CCATATTTAA TGGAGGATTA TTTGCACAAG ATAAGGTTAA 180

ATATTTAAAT AATGAAAGTT TACTCAGTAT TAGTGAGATT GAAGAAATAT TAGTCAAAAT 240

ACTTTTCTTT GAAGAAAAAA ATATTAAAGA TAAAAAATTT GTAAAATATT CAAGGCTAGA 300

TCCTAAAAGC TTTGGAGAAT TATACGAAAC TCTACTTGAA TATGACCTAA GAATTGCAGA 360

TACTACTGTT CATCGTATTG TTGAAGACGG GATTTATCTC ATTCGTACTG AAGAAGAGCT 420

TGAAAACAAT AAAGTAAACA AAATTGCTAC ATATCTTAAA GGGAATATTT ATCTTACATC 480

TAGATCACTT GATAGAAAGA AAAGTGGGGC ATATTATACT CCAGATGATT TAACTGATTT 540

TATGGTTATA TCATCAATTG AAGAGCAGCT TAAAACCAAG TCCCCTTTAG ATATAAAAAT 600

CATTGATAAT TCTTGTGGAT CAGGGCATTT TTTAATTTCT TGTCTAGATT ACTTAACAGA 660

AAAGGTATGG TACGAGCTAG ATAAATTTGA AGATGTAAAA AAAGAGCTTG ATAAAGAATA 720

TGGGATTATT CTTAAAGAAA GTGAGGAGTA TGATATTCAA GATAGTATAA GTAAAGAATT 780

GGTGCTTAAA AGGATGCTGC TAAAGAGGTG TATTTATGGT GTTGATATTA ATCCTATTTC 840

GGTTGAAATT ACTATGCTAA GTTTGTGGAT TAATACCTTT ATTTTTGGAA CGCCACTAAG 900

CTTTATTGAG CATCATATAA AAACAGGAAA TGCTCTCTTG GGATAT 946

(2) INFORMATION FOR SEQ ID NO: 101:

(i) SEQUENCE CHARACTERISTICS:
(A) LENGTH: 913 base pairs
(B) TYPE: nucleic acid
(C) STRANDEDNESS: double
(D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 101:

CTTTGACTCA AAACTTTACC CTTTAAATTG CTAACTTTAA CTTGAAAATA CTAAACTTTA 60

ACCCGAAATA ATAAAACTTT AACTAGAATT TTTCAAAATG ATAAAACTTT AATTTTTGCA 120

ATTTTATTCT CTTGTTTTTT TTAAAACGAT TAGAATAATC GTTGAkCAGG TTTATTGATT 180

ATCAATAAAC CTGATCTATA ATATTATAAG CGGTTTTTGC AAGTTTAATA GGAGCTATAA 240

TATCCATGAA CAAATTATTG ATATTCATTA TTTTATTAGT CTTTTCATGT AATTTAAGTA 300

ATTCTGATCA AAATAATCCA CTAAACATGT CAAATAAAGA AAAAATAAGC GAATATCAAA 360

TAAATGAGTC GTCAAACAAA TATTCAATTT TCAAACGAAA TTCAAGCGTT AAAAGATACA 420

CGTTCAACCA TTATTACTAA CCAAAATGAT AATATTAATT CTACTATTAA CTACCCACCT 480

TATATTCAAA CTATCTTAAA AATAGAAAAA CAAGTTGACG GAAATATTAT TATTAATGGG 540

ATGACTAAAG AAAGTGGCAC AGAAACTAAA AAGCTTTTAG AAATTCCAAA TGGGAATATT 600

TCTCGACTTA AAGATGCAAT TCAATATGGA GGAAGTTTTA GGGCTAAAGA TGTTAGAGAA 660

AATCAAACCC AAAAAGAAAA CAACAAAGAC TCGCATATTC ATGTCGACrA TTTTAAAGAA 720

TACATACATT TAATCATGCC TAGCATTAAC AATAATGCTG ATAGTAGTAG TAGTTATTAC 780

TATACCAACT ACATAATAAA TGGAGACAAT TTGTTAAGAA TTATTAGCAA CTTATAAAAr 840

ATCTTTATAA ATTACCAATA TTCTTGArAA TTTTAATACT ATTTkgTTaT ATACTATAAT 900

ATTAAGAGAA AGA 913

(2) INFORMATION FOR SEQ ID NO: 102:

(i) SEQUENCE CHARACTERISTICS:
(A) LENGTH: 910 base pairs
(B) TYPE: nucleic acid
(C) STRANDEDNESS: double
(D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 102:

TCGCTATnnG AGCTCGGTAC CCTGATAAGG CGAGTGTGAC GGGGATTGCT AAGGGAATAA 60

AGGAGATTGT TGAAGCTGCT GGGGGGAGTG AAAAGCTGAA AGTTGCTGCT GCTGAAGGGG 120

AGAATAATGA AAAGGCAGGG AAGTTGTTTG GGAAGGCTGG TGCTGGTAAT GCTGGGGACA 180

GTGAGGCTGC TAGCAAGGCG GCTGGTGCTG TTAGTGCTGT TAGTGGGGAG CAGATATTAA 240

GTGCGATTGT TAAGGCTGCT GGTGAGGCTG CGCAGGATGG AGAGAAGCCT GGGGAGGCTA 300

AAAATCCGAT TGCTGCTGCT ATTGGGAAGG GTAATGAGGA TGGTGCGGAG TTTAAGGATG 360

AGATGAAGAA GGATGATCAG ATTGCTGCTG CTATTGCTTT GAGGGGGATG GCTAAGGATG 420

GAAAGTTTGC TGTGAAGAAT GATGAGAAAG GGAAGGCTGA GGGGGCTATT AAGGGAGCTG 480

GCGAGTTGTT GGATAAGCTG GTAAAAGCTG TAAAGACAGC TGAGGGGGCT TCAAGTGGTA 540

CTGCTGCAAT TGGAGAAGTT GTGGCTGATG ATAATGCTGC GAAGGTTGCT GATAAGGCGA 600

GTGTGAAGGG GATTGCTAAG GGGATAAAGG AGATTGTTGA AGCTGCTGGG GGGAGTAAAA 660

AGCTGAAAGT TGCTGCTGCT AAAGAGGGCA ATGAAAAGGC AGGGAAGTTG TTTGGGAAAG 720

TTGATGCTGC TCATGCTGGG GACAGTGAGG CTGCTAGCAA GGCGGCTGGT GCTGTTAGTG 780

CTGTTAGTGG GGAgCAGATA TTAAGTGCGA TTGTTAAGGC TGCTGGTGCG GCTGCTGGTG 840

ATCAGGAGGG AAAGAAGCCT GGGGATGCTA AAAATCCGAT TGCTGCTGCT ATTGGGAAGG 900

GTGATGCGGA 910

(2) INFORMATION FOR SEQ ID NO: 103:

(i) SEQUENCE CHARACTERISTICS:
(A) LENGTH: 888 base pairs
(B) TYPE: nucleic acid
(C) STRANDEDNESS: double
(D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 103:

AACATGTAAA AGAATAAGCA TTAACTCGCG CATTCTTTGA TTTAAAACAA CCACCCGAAC 60

TACTAAAAAC CTTATTTTCA ATCGAACTCA TTGATTTTGA ATATTTTTTA AATTTTAAAA 120

GAACATCGTC AAGTTCTTTA ACTGAATCTA AATAAGGATC TTTTGCCTGT ACTTCTTCAG 180

CCTGTCTTGT TTGACGTTTA GATCTAGGAG CAACTGGAAT TTCTGATTCT AGCCCTAATT 240

GTGGATTATC ATCAACATTA GGAGCTTTAG CTTGCCCTTT GCCTTTTAAA GCCATAATTT 300

AATTACCTTT TAAAGCTCTA TTCCCAAAAA CACTAGCAAG CACTATAGAT AACTCTTCGG 360

TTAATTTATG TACTTTTGAA AGTGCTATAG CATTAACAGA TTTATCATTT CCCCCATTCT 420

TTTCAAGCTC TCCTTGTGCA TTAAAATGCA GCTTATCACC TGGGTTTACA CCATTTCCAT 480

TTTTCTTAAA TGTTAAATAC CCCGTGAAGT TATTTGTAAT TGGAACTACA GTTGCCATGC 540

CAGTAAACTC ATCTATATCA GTGCATATTC CGTACAAGTC ATCCCCACCA CCAGCCTCAA 600

CTTCTAGTTC TGTTGTACCA TCTGCACTAA AACTAAGCTT GACTCCACGC TTGTATGGAT 660

ACCCTTTAGC AGGATAATTT TCTATTTTGT CTTTACTACT AGTAAAAACT CCATCCGAAT 720

TGGAGTAAAT TAGATTTTTA TCTCTAAAAT CTACAGAATT ACTAAGCAAA CCAGTATCTT 780

GCTGAGGATT TTTCATTAAT GCTTTAATTT CTGCAACTTT TTTATCAAAC TCTTCTTTTA 840

TTTTTGTAAT ACCATCGCTC ATTAAAAACT CCTTTAAGCA ATACTGGT 888

(2) INFORMATION FOR SEQ ID NO: 104:

(i) SEQUENCE CHARACTERISTICS:
(A) LENGTH: 883 base pairs
(B) TYPE: nucleic acid
(C) STRANDEDNESS: double
(D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 104:

GATAGTAATT AAGTCCTACA ACCAATAGTG TTACTTTGCA TTAATTTGTT CTTTCCTTAT 60

CGATAGGTCT CTTCCTTCCT GATTGAATTT CAGATCATTA GATATTTTGA GACTTTCTTC 120

ATCAGAATTA ACTCAAGTCA ATGCATTGAT TGATTTTCTC ATTTAATGGA GCTAGTGCTT 180

TATTTATTGC TGGGGTTAAT GCACTCTCAA GTCTTTCCAT ATTTGCTGTA TAGATTAATT 240

TmTAATGAGA ATACAGCTCA TAAACCAAAA AGAATCCTTT ATGTGCAATT TCATCAAATT 300

CATCTTCAAA TTTAGAAAAT ATATCAATAA GGGTTGATAA AGACGTAAGT CCAAGCTCAA 360

CATTATCTTT GGATAATTTC ATAAGTTAAT CTCTTTTTTT AATGTGATTT TTGCCATTAC 420

CATTGCCATT CTTAAAAATC TTGCCTATTA CAATAGTCAA TATGTCTTTT AATAAAGGCT 480

TGAGAAGAAT TAACACTCCT AAAACCAATA CTGTTACAAA AATCATTACG GCTATAAGTT 540

TAATTTCATT TAAATTGATA AGAAGTTCTG TTAATTTAAT AGTATCCATT TTTTAATCCT 600

TTATTTTAAT TTTTTATTTG TATATACATT ATATCAAAAT CGTAATTTTT GCTAAAAAAG 660

TTTGCAGCTT TTAAAGCTGC GGGATGGGGC CCCCTGATAG GTAGGCTCTT TTTTGAATAT 720

ACCATCCTTT ATACATGGGA AATCTACTAG ATAGTCCTTG GGGGAGCGTC TGrTTGCTCA 780

TAAGCCATAC TTGTTTCACT TTCATCGGAA TATCTTAGAT AAAGTACTTT ACTCTCGCTA 840

TTACTGTAGT GTTCTGCGTC AAGCTCAATA TCAAGGTAAA TGG 883

(2) INFORMATION FOR SEQ ID NO: 105:

(i) SEQUENCE CHARACTERISTICS:
(A) LENGTH: 857 base pairs
(B) TYPE: nucleic acid
(C) STRANDEDNESS: double
(D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 105:

CTATTAATTG CCAAAAAACT AAACCAAACA ACACTTAAAA GGAGAGTCTT ATTGTGATAA 60

TGAGATATAA AATGAAAATT TTAACTAAAA ATAAAACTTA TGAATATCCG CTGAGAGTAC 120

TTCCCGTCTA TGAATGGGAT AAAGTGCTAG GATTTAATCA AAGTGACGCT GTTTTAAAGC 180

TTAATGAGGT TAAATTCTTA AGAGAAATCA CAAGCTTAAT GATAAGTCCA AAATTTTTAG 240

ACGAATTCTA TGTGATTTTG GATCAAAATA GAGAATTTAT TTCTTATTAT AAGGACTATC 300

TTGTTGCAAT AATTTACACT GCACAATTTA ATACTTTTCA TTTAGACAAT AATCTAAAAA 360

AGCCCGCTTT AGTATATTTG AGTGAGTATG AAAATAATGT TGGTGATTTT GTTGCTTTtG 420

ACTATATTAA TGAAAATTTT GATTATGAAA AAGTAGCCAC TTCGCTTTCA TCAATTACAT 480

CAAATTCCAA TGAGCTGGTT GCTAAATGAG CAAAAGAAAT AGAGATATTG ATAAAGCTAT 540

TGCAAGTCTT GATGAGACTA GAAAAAAATA TTTTAACTTG CTTGACGAGA TTAAGAACGA 600

TAAATACTTT TTCCCAGTAA TTATGAATAT TTGCTCATAC TACTCGGTTA AAAAATTGCC 660

TTATGACGAG CTTTTAGAAG TCAATAGACT TGCTGAGATT AAATTAGAAA AAGAATTGTA 720

TGAATTAATT TTAAGCAAGT GAGGACTTAG TGAGCGACAA ATTCACCATT AAAtTTAAAG 780

GtATTCyTGA TCaTGCTGCA ACAAAAAAGG GCCATTGAAC CAGGATATTT CTAAAATGGn 840

AAAATTATCn TAAACCT 857

(2) INFORMATION FOR SEQ ID NO: 106:

(i) SEQUENCE CHARACTERISTICS:
(A) LENGTH: 846 base pairs
(B) TYPE: nucleic acid
(C) STRANDEDNESS: double
(D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 106:

AGTTGTTCTT TTGCGAGATG CGCGTCAAAA CGTAGCATAG TATTGCCAAA AACAGTCATT 60

TTAAGTAAAG AAAGCATATC CTTATATAAG GATGTTAGTA TTCCACCGTT TAATGATATG 120

TTTTCACCAG TCATTACCGG ATTGTAGCTT ACATATTCCG CTTTTCTATC ATAATAGTTG 180

ATAACTGGTC TTTTAGAACA ATTAGTATTG TAAGTGCGTG TTATGAGTTC ATTTTTTGGT 240

TTTATAAAAA ACAATTGAGG AACATATCCA AAACCTTTTA GATCCATTCT AGGAAATAAC 300

ACTAAAAAAT TATCTGCTCC GAAAAGGGCA AATATTTGGG TTATTACATC TCTTATTATT 360

CGAGTAATTT CCCCGATTTC TTTCTTTTCA ATATCATTAA TTTTTTCCTC GATTTTTTCT 420

TTTCAATATC ATTAATTTTT TCCTCGATTT TTTTCTTTTC AATATCATTA ATTTTTTCCT 480

CGATTTTTTT CTTTTCAATA TCATGATTGT TAGTAATTTT ATTATTAATA TCTATTTTGT 540

TAGCTGCATT GTTAGCAATT TTTTTGTTAC TTGTCATAAG TAATTACCTT TTGTAAAAAT 600

TATGGTGTGC TGTTAGCATT GTCTTGATTT TGAATTTCTT CTTGTAGTTT TTTTAGAGCC 660

GCACCCTCAT CTCCGCCCAT CCATCCAGGT AGCATCGATT TTAATTTTGC AAAGAAATAA 720

TTAAGATTAA AAATACTTTT AATGCCATaA TTATGGGATT AATAAGTGTG TTTCAAACGC 780

AAAGTCTTAA AGTAATAGTT ATCTTATTAA TGAGGTCAGT AAGGGTCCAA GACGTAGTGG 840

TTAAGT 846

(2) INFORMATION FOR SEQ ID NO: 107:

(i) SEQUENCE CHARACTERISTICS:
(A) LENGTH: 840 base pairs
(B) TYPE: nucleic acid
(C) STRANDEDNESS: double
(D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 107:

CAATGAAATA GTTAGAGAAA TTAAAAATGT TATTAAAAAG CACAATTTGG AGCTTGATAT 60

TGAGCAATAT CCAATTTCTA TAGAGGGTCA ATATGGCATA GTTGATTATA TTAGGACTAC 120

ATTCTACAGT ACAAGTACTG GATATGAATT TTCTTTTGAT ACGCGAATTC CTACAGAARa 180

TTtACAATGG aACAATGAAA ATGGGTCTAA AGTTACAAAT ACAGTGTATC AGATGTTTGG 240

TTCAGGCATT ACTTATGTCA AAAGGTATGC TTTAGTTGCA GCTCTTGGTA TAGAAAGTGA 300

AATAGATACT GATGCAGCTC CTATTTACAA TAACCACGAA AACGAAAATT CTATGCCTAG 360

CAAGCAAGTT AGTGTTAATC AAAAGCAAGA ACAAAAAAGA GAACAAAAAC AAGAAAAAAA 420

TCAACTAAAC AACTTTAATA AAAACTTAAA ATCTGGCAAG GCTTATTGCT ATGAAATTTT 480

TAGAGACGCA CTGTTTAATA TAAAAAATTG GGTAAATGAA GGTGAAGAAA AAAATAATAT 540

AAATGCTCTT ATTCGGGCAT TATGTACTGA TAATGATGAT GCTTTAGAGG ATCTTTTTGA 600

AAAGAATGCT GAGCTTAAGA ATATAGAATA TTGGGTAAAT ATTCTAAAAA AATATTTCAA 660

TAAAACCnAT AGATTTGATG ATCTAAATAA GCTnAAAGTT TTTATGTCTG ATAATCGGGA 720

TGTTTATAAA ACAAAAAnTA TTAAAATTCT TTTGCATGTT AAAAAAAAAA AAGACAATTT 780

AATTATATnT TTGCCAGTGT TGCCATATTA AAGCCCCCCA ATAAGGGGGC TGTTTAnATT 840

(2) INFORMATION FOR SEQ ID NO: 108:

(i) SEQUENCE CHARACTERISTICS:
(A) LENGTH: 814 base pairs
(B) TYPE: nucleic acid
(C) STRANDEDNESS: double
(D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 108:

GGCCGACTTT GTTTTATCTA CAATAACTTA TTTTAGTTTG AAAATTACAA GCAAGCCTGA 60

AACCATTCAT TATCCCTATG GACACAAACT AATGGAAAGC TTAATAGCTT TTATCATGGG 120

AATAATTATA CTTATGACAG GATTTACACT ATTTCTAAAT ACAACCGGAT TAAATAAATT 180

TATCACTCTT GGGGGAGAGT CTGGATTTAA TCTACACATA CACCAGAACA AAAATAAAAA 240

TGATACTATA TATGAACATG ACCATTGCCA TTCACACGAT CACGATCACG ATCATAACCA 300

CGACCACAAC GAAGAAGACA AAAAAAACAT ACTAGAAATA TTTTCAAATA AATGTCTAGA 360

AGCAAAAGCA AGCTTTCGAT AAAACCCGAA GTTGTTTCGC TAAAGTGACA AGGATTAAAC 420

AGGATTGTAT TTTTCAGCAG CCTATTTTAT AAACGATCTG CATTTAGTAA ATAGTTTTTA 480

GTTAGGAAAT AATGTAGGAT TACTAAGTGT GATGTCTGAG AGAAGGGACA AGTATTGTAG 540

CGAGCTTAAA TCCTTATTAT CGTTGGCCAG TAATTTAGAG GTAGGGGATC GGGATAAAGG 600

ATTGGCCAGT TTATAAGTTG GAGGGAAGGG CAAAGGATGC CTTAAAATCG GTAATCGCTC 660

CTTAAGGTTT AGGGTTAACA AGTTTGGCCA CCAATTAACC TCCAAAAAAA GGCCAGGCAA 720

AAATACCCAT TAAAGGCAGG ATTTCCGGTT TGGGAATTGG CCCCGGACCA CCTCCAAAAG 780

GGCCATGGAA GGTTTGGGAA ATTGGTTAGG TAAA 814

(2) INFORMATION FOR SEQ ID NO: 109:

(i) SEQUENCE CHARACTERISTICS:
(A) LENGTH: 808 base pairs
(B) TYPE: nucleic acid
(C) STRANDEDNESS: double
(D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 109:

GGCAAAACCA GTTCTATTTT GTTCAAAATC AATATTCGAC TCTTCAGAAT TTAGTTTTAA 60

ACTTTTGATT TCATATTTGT TTTCAAACTC TTCAGTTGAT TCAAATGCTA TTATCTTAGC 120

TATAGGTACT TCTTGACCGA ATATTTTATA GTTATTGTCA TTAATGCTAT AATTAAGTAT 180

AGCTAAAGGT AAGCAATGAA TAAGTTTTGA ATCGGATGGG TGAAAAAATA TAAGAATACT 240

TAAGCTCTCT AAGATTTTCA ATTAAATTGT TATTTTTATC TTTTATATCT TTTAAATCAC 300

CAGATTTTGC CCATGTAACA TAACCACTTG CAAGTATTTC AAATGCCTCT CCTTCTTTTA 360

TTTCGTTTAA TTCGGGTTCC TTAAAGGTAA GACAAATATT GCTATTACTC TCTTCAATGG 420

GCTTTTTATC ATATTTTATA ATATTTCCCT CTTCAATTAT CTTAAAATCC AAAACCTCGG 480

TTTTGATACT GTTGATTTTT GAAGATTTTG GTGTAGTCAA AAAGCATGAA TAAAATAATG 540

GTATATATAA AAATATATTC AAAATACATA TGTTCTTTTT CATAAAATTT TTCCATTAAT 600

TGTATTTCTT TCTATTTCTC TCTTCACTTT TAAATATTGT TGATAAGCAG TGGGTCTAGG 660

CATAAAACGA TCATACTCAG GGCTCCCCTC TTCGCCAGAA TACTTAATAT CTGGAGAATA 720

TAACTCGCTT ATACATGAAT ACAGCCAATA AACTTCACTT TTAAATTTAT TATTCTGCTC 780

ATTTTTTACC CTACCAAACA ACTTAATC 808

(2) INFORMATION FOR SEQ ID NO: 110:

(i) SEQUENCE CHARACTERISTICS:
(A) LENGTH: 804 base pairs
(B) TYPE: nucleic acid
(C) STRANDEDNESS: double
(D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 110:

TGGGAAACAT TATATCTAAC AACCCCTAGT GGTACTTTAC TTGAGGGGGA CATAGAAATT 60

GATGGCCTCA ATTCAACTGG ACAACGAAAA TCCTACAAAA TATCGCTAGG AAAAAGAAAA 120

TATGTTTATA TGAAAGTAAA GTATAAACTT GACCTTAAAA ACTATCTCTA CTTAAACATA 180

GACTCTCAAA TTAGAGACAT TTATTCTAGG ATTATTTCAA ATAACTATTC TGATATGGGA 240

ATTAGCTTTG AATATCAAGA CTTTTTTGCT CCAGTTAATG AAGTTAAAGG AATTAAATTT 300

ATGGAAATAA GTGCCTGTAT TAAAGACACA GACACTGAGA GTATTGCAAA AATTACTGAT 360

AGCGATTTTA AAAAAAATCA AGATATTACT ATTACTGATG ATACAATGCT CCTTTTCAAT 420

ACTACAGATA GATTGCTTAT TGATATTGAT AGTTAACAAA TATGAAAATA CCTAATTTAT 480

TCAATGGCAC TGAAGTTCAT AAATTTATAC TTACAGAAAC AGAATATGCA CAAGCATTGC 540

TTAATGAACT CAAGTCTCTT AATTCTAACT TCCTATCCAT TAATGTAATA GAAAATATAA 600

AATCAAGATA TATTGCAATA TGGATATCTC AAGTTTTATC TATCTTTTAT GCAAAAACTC 660

AAACTTTACA AAGTATTACA AGCAATATTA ATAGCGTTAT TTTTGCTTTA CGCCATATTG 720

GTACTGATGA GTCGTTTAGA CTAATTTTCA aGGCCTTTTa AATGTGGACA TGAAGTTACT 780

ACTCCTGAAG CGGGGGTATG GAAA 804

(2) INFORMATION FOR SEQ ID NO: 111:

(i) SEQUENCE CHARACTERISTICS:
(A) LENGTH: 800 base pairs
(B) TYPE: nucleic acid
(C) STRANDEDNESS: double
(D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 111:

GCGTGAGATG TTAATTTTTA CCnGCTTTAA AGCAGAATAG TCCATCCCCA TGAGGAGCAT 60

AGCTTAAACT nCCCCTTAAG GTTTTTAAAA ATATCAAAAA CCTTnGTTTT TGTTCATCAA 120

TTCCATCAAC AGTGATAAAC GTTATTAAAT ACCAATTCAA AATTATTGCC TAATCTTAAA 180

yCCATACTAG TAATTATACC ATATTTAGCA AAAATTACTT AAACAAACTA TTAATATCAG 240

AATTAATTTG AGCGGTTGCT ATTTTTAAGC TTGATTCGTC AATTACTGAG TCCCCTATAA 300

TTTTTATACC ATTGATAGCA CTAACAATAT TATCTAGAAT TTTTTTTAAG CTAGTTGTTT 360

GGTTTGCTAT TTCAATTTTA TTATTCGCTC TAATTTTAAC AGTATCAGAG ATTAGATTTA 420

AAGTCTTTGG GCTAATTGCA CTAAGTATAT AAAAATGATG TTTGTCAAAG TGAATATCGT 480

TATTTTTATC AAAAATATTA ATGCTTGATT GAAGTAGTAA AACGCAATCA CCTTTTGATA 540

GTTCTAAACT GATATTAGAG ATATTTTTTG TGTGAATTTC TAAATCTTCA AATTCGGGTA 600

TTGTAACAAT AGCTTCTTGA GTTTGATGTT TAAACTCCTT TACAGTGCCA ATTTTAATTA 660

TAAAAATGTT TGAATAAATC CAATTTTTAA GGTCTTCTTG AGCCAATGCC TGGCCATAAA 720

GGCGTTGATT CATTCtGTAA ATTTCATAGT CTTCaTTCaT TCtAATTCyA GTCCCCTTTA 780

tTTTTTACgt TTTGTATTAG 800

(2) INFORMATION FOR SEQ ID NO: 112:

(i) *SEQUENCE CHARACTERISTICS:*
(A) LENGTH: 798 base pairs
(B) TYPE: nucleic acid
(C) *STRANDEDNESS*: double
(D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 112:

GAAATAGCTT TTTAAGTTTT CTAAATCATC TTTAATATCA ATAACTTCTA TAAAGTATTA 60

ATGATGAATT TTTGGGAATG TTTTATGGAT TTAAAAGATT AACAAGACCA CTTTTTTTAA 120

ATACGAAGAT ATTATTACAA AAACTATCAA AACTGTACCC ATGTATAAAA TTCATTACAT 180

AGAATTTAGA TTTAAGAAAG GAAGTGTTTT TTGTTATATA AAAGCAATTC ATGTTTTAAT 240

AAAAAAAGAA AAATTTAAAA AAAATATGCT CAAAGTCTAT TAGAGAGAAT AATTAATCTA 300

GAACATAAAG TATTAAAGAT AAAGCAATTT TTTTAAAAAA TATATAAAAA TCGAAACAAA 360

AAATTAAAGA TATAGTAAAA TTGTATTTGT AGCAATATAC TTGTGCTAGA GGCTATGAAT 420

CTCTAAAGAT TTTAGCAGGG GAGAAAATAT GAAAAAAAGT TTTTTATCAA TATACATGTT 480

AATTTCAATA AGTTTATTAT CATGTGATGT TAGTAGATTA AATCAGAGAA ATATTAATGA 540

GCTTAAAATT TTTGTTGAAA AGGCCAAGTA TTATTCTATA AAATTAGACG CTATTTATAA 600

CGAATGTACA GGAGCATATA ATGATATTAT GACTTATTCG GAAGGTACAT TTTCTGATCA 660

AAGTAAGGTT AATCAAGCTA TATCTATATT TAAAAAAGAC AATAAAATTG TTAATAAGTT 720

TAAGGAGCTT GAAAAGATTA TAGAAGAATA CAAACCTATG TTTTTAAGTA AATTAATTGA 780

TGATTTTGCG GGATCCGT 798

(2) INFORMATION FOR SEQ ID NO: 113:

(i) SEQUENCE CHARACTERISTICS:
(A) LENGTH: 798 base pairs
(B) TYPE: nucleic acid
(C) STRANDEDNESS: double
(D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 113:

AGCTTTTGCA TAATAATTTT CATCAAATAA TTCCCATATT AAATCCTCCC AAATATCATT 60

AATTTTTACT TACAGCTTTA TTTCCAAATA CTGCTACTTT TATTAAATAA ACATCGTTAC 120

TAATTTGTTT TGCATCAGAC AACGCTATTG CATTAATAGT TGCCTTATTT GGTGGTGCTC 180

CAGTCACCTT TTCAAGAGCA CCGTCTTTAT TAAAAACAAG TTTGTCTTTT ACTTTAAGCG 240

TAGAATCTTT TGCTACTAAA TAACCCTCAA AATTATTTGT AATCGGAACA ATAGTGGCTG 300

TTTTGCTAAA CTCATCTATA TCAATGCATA TTCCGTATAA ATCATCTTCA CCACCAGCCT 360

CAACGTGGGG TTCATAGTGA ATTTGATCAG CTTTTTCCTC TTGAATAACT CTTTTTACCC 420

CACGCTTATA TGGATACCCA GAAAATGGAT GATTTTCTAA TTTGTCAAAT TTGCTGGTTC 480

TAGTGCCTCC AGAGGCAAAA AATTGTATGT TTTTATCTCT AAACTCTACA GAATTGCTAA 540

GCAAACCAGC GTCATGCTGG GGATTTTTCA TAAACTTTTC AAGTTTACTT CTCTTCTCTT 600

GaTAATCTTT TACTAATTGC GTTGTGTCTG cCATTTGTTT AACTCCTTTT ATTGcCCAAG 660

GgCkAwCCrC CAGCTtCAGG TGTTACTGTT TTCTCAAGGG CCTCTATTGG CCAAAAATTG 720

GCAAACTTTT TTTTTAAATT CCCAAAAAAA AATTTTTAAA AATTTAAAGG GAAAAATTTA 780

AACCCTTTCC CCTTTTTG 798

(2) INFORMATION FOR SEQ ID NO: 114:

(i) SEQUENCE CHARACTERISTICS:
(A) LENGTH: 783 base pairs
(B) TYPE: nucleic acid
(C) STRANDEDNESS: double
(D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 114:

GGACTCAAAA CTTTACCCTT TAAATTACTA AATTTAACTT GAAAATACTA AACTTTAACC 60

CAAAATAATA AAACTTTAAC TTGAATTTTT CAAAATTACA AAACTTTAAC CCAAAATGAT 120

AAAACTTTAA TTTTTTGTAA TTTTTACATA AAAGTGTTAA CTTTAAAATC CCAAACTTTA 180

TAATTTTGGG AAATTATCAA TACTTTTTTA ATTTATTCTT TATTTTCAAA ATAATCTTTA 240

TATACTTATA TATTATGTAT AAGGCTATAA AAGAACAACA AGAAATAGAA ATAGATCATG 300

CATGCAGAAT ACTTATTCTT ACCGCAACAA TATTTGAAAT AAATTCAATA TTCGAAAATT 360

ATTATCAAAA AACTCTACTC AAAAAGTATA ACGAAAATCT CAAAAACAAA AATCTACCTC 420

CTAGTAATAT ATCAACAATG AAAAAATACT TAAATCAATT AGAAAAAGAA ATAAAAATCA 480

TAGCAAAATT CTATTTTAAA AACGATCAAT CTCTAATTTA TTGCAAACTT AATTATACCC 540

TAGAAAAAAT TTGTTTAAAA CTAATAAAAT TCTACAAAAA ATTCTACAAA GAATTAAAAC 600

AATTTACACA AAAGAACATT ACTACTTAAT TGTAAATACA TTATAAAATA ATCTTATGCA 660

AATATTTAGA AATACAAATT GTAAAGATAT ATATTTTTAT TTAAATAAAT AATAAAAATT 720

GCTGGCACAC TAATTTGGAA AAATCTTTAA AAGAnATACT AGGTATGAAT AGCnAAAATA 780

AGC 783

(2) INFORMATION FOR SEQ ID NO: 115:

(i) SEQUENCE CHARACTERISTICS:
(A) LENGTH: 768 base pairs
(B) TYPE: nucleic acid
(C) STRANDEDNESS: double
(D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 115:

TTCTATATAA ATATTTTGTA ACTTTTTTGC TTATTACAGA CTAAGCCTAA ACGTCCTACA 60

ACCCCATAAA TGCAACGCTC TGCAGCTTGA CACATTTAAA GTTTGGGCTA CTCCCTTTTC 120

GCTCGCCACT ACTAAGGGAA TCTCTTTGAT TTCTTTTCCT CAGGGTACTT AGATGGTTCA 180

CTTCCCCTGG TATCGCCTCT ATTATTTAAA TAATAGATAG CTAGCATCTT GCTAGCTGGA 240

TTACTCCATT CGGTAATCTT GGGATCAATA AATGTTTGCT TCTCCCCCAA GCTTTTCGCA 300

GCTTACCACG ACCTTCTTCG CCTTAAAGCT CCTAGGCATT CACCATAGAC TCTTATTACT 360

TTGACCATAT TTTTATCTTC CATCTCTATT TTGCCAATTT ATTTATACAA CATAAAATAA 420

TATATATCTT TGTTTAATAC ATGTCAATAT ATATTTTATT TTTTATGTTA TTTAAACAAC 480

ACATTCAAAA ACACCAATAT TTAAAAAACA TAAAAATAAA ATCAAAGTTT AAAGTATAAA 540

AATAAAAACC CTGGCAATAA CCTACTCTCC CGCGAACTCG CAGTACCATC AGCGAATAAG 600

AGCTTAACTT CTGTGTTCGG AATGATAACA GGTGTTTCCT CTTTTCTTTA ACCACCAGGG 660

TTTTTACAAG GAAGACAAAA ATATgGcCAA AGATACGGGT AATTAGTATT AGTCAGCTTA 720

ATATATTGCT ATACTTACAC TTCTAACCTA TCGACCTGGT ATTCTTTC 768

(2) INFORMATION FOR SEQ ID NO: 116:

(i) SEQUENCE CHARACTERISTICS:
(A) LENGTH: 765 base pairs
(B) TYPE: nucleic acid
(C) STRANDEDNESS: double
(D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 116:

CTTTACGCCT AACTTACCCT CCACGTGTAC AACCCCTTAA CAACCCCTTA ACTTACCAGT 60

GACCCCCTTA ATATGGTAGT TATGGGGGAA CGCTTAGAAT AAAAAAGTCA TCTACGACAC 120

CCCCGATCAT AGACCTGACT CTTGTTATCC CAAATCACTT CAGCGCCCTC GCAACTTATG 180

GGAAAAAGTT CCTAGAAAGA TGTATAGAGA AGTGGAATCA AAGTAATAGG CAATTCGCAA 240

GTGAATAAGG GAGAAAGGAT TTCCTATGTT ATAGGGAGAC GCTAGACATA GTGCTTGCGA 300

GACTGGATTG TGCTTGATGG ATAGAACCTA GTTTAGTGTG TACATCCAAA AAATGGACTA 360

AATCAATAGT ATAAGGCGAA TTGCCAGCGA TGAGGTCTGA ACCAAATTGC CTCACATCAG 420

GCGATAACAG TTACTACTAA CAGTTGCCAC TTCGGCTACT CTATCTTGCG TGCTTATTGT 480

AGCACTCTGG AGGTCCTGTT GTTAAGCCAG CATTAGCACA GCTCCTCCAC TGCGGGTTGC 540

GAGTATAGAG TAGTCCTAAC TGGCAAGGAT TCCCCCTCTG GTTGCTAGAG GTCGAATTAC 600

CCACCCAACA ATAGTTGCAT TGTTGGGGGG GTGGGTACCT ACTACTCGGC ATATACTCCC 660

CCCCTTTCGA GACCTCCCTC GAGGGTCGAG GGAGCATTTG ATCATAGACG TTCATCCCAG 720

ACATGGCCTT TCGGGTTTGA CGTCTCGCGA CCCCCCTTCG GGGAC 765

(2) INFORMATION FOR SEQ ID NO: 117:

(i) SEQUENCE CHARACTERISTICS:
(A) LENGTH: 755 base pairs
(B) TYPE: nucleic acid
(C) STRANDEDNESS: double
(D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 117:

GAGCCATTAT TGGCTCCTAT TTTACTAGCT GCACTTGAAA TTCTTTTCTT TTTTAGGAAC 60

AAATAACGCT CTTTAAATTA AAAGGCATAA TGCTATATTG TATTCTAAAT CATATACAAA 120

GGACAGTTCT TTATATCATA AGTGCAAAAA TAAAGTCATA AATTCAATAA AAAGGAGGAA 180

AACTCTTCTA GAGTAGTAGA AGAGCAACCA AAAATTAATG AAAATTTTTC TACACAAGAA 240

TCTATACAAA AACTGCCCCT TTACTGCAAC ATACAAAACG TGAATCTTGT ATATTACAAT 300

AATAGATAAT ATTATTGCAA CAATCCTAAA TTACAAATAC AGAATATGTT ATTAGCCCCA 360

AAAAGGGGCT AATACATTTA CTTTAAATTA CAAGTTATTC GAACCATAAT TGTTCAATAT 420

TAATTTCAAA TCTTTTCTTA TAGCAAGAAA TTTTTCATAA ATCAATATTA GATAATCATC 480

AAAATTGCTT TTATCAAGCA CATACAAAAG TTTAAAAAAA TCTACATCAT CAAGACATAA 540

ATAGAATATG AAAACCTTAT TTTCAAACAC ATTATCACCC AGCTTTACTT TAATTTTACG 600

AAAAAGGTTG ATTAATTCTT TAGACTTTTT TGGCCCCAAA TTAAAAAAAA ATTCATTTAA 660

AATGTTTTGA CTTTTAGGCG GAGACAATAT ATTTATTGTC TCCGCATCAT TTTCTATATC 720

TAAGAATCGA CTCATAGGAA CTTATAAATG ACTTT 755

(2) INFORMATION FOR SEQ ID NO: 118:

(i) SEQUENCE CHARACTERISTICS:
(A) LENGTH: 753 base pairs
(B) TYPE: nucleic acid
(C) STRANDEDNESS: double
(D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 118:

CTCAGCCAAG ATACTGGTTT GCTTAGTAAT TCTGTAGATT TTAGAGATnA AAATCTAATT 60

TACTCCAATT CGGATGGAGT TTTTACTAGT AGTAAAGACA AAATAGAAAA TTATCCTGCT 120

AAAGGGTATC CATACAAGCG TGGAGTCAAG CTTAGTTTTA GTGCAGATGG TACAACAGAA 180

CTAGAAGTTG AGGCTGGTGG TGGGGATGAC TTGTACGGAA TATGCACTGA TATAGaTGAG 240

TTTACTGGCA TGGCAACTGT AGTTCCAATT ACAAATAACT TCACGGGGTA TTTAACATTT 300

AAGAAAAATG GAAATGGTGT AAACCCAGGT GATAAGCTGC ATTTTAATGC ACAAGGAGAG 360

CTTGAAAAGA ATGGGGGAAA TGATAAATCT GTTAATGCTA TAGCACTTTC AAAAGTACAT 420

AAATTAACCG AAGAGTTATC TATAGTGCTT GCTAGTGTTT TTGGGAATAG AGCTTTAAAA 480

GGTAATTAAA TTATGGCTTT AAAAGGCAAA GGGCAAGCTA AAGCTCCTAA TGTTGATGAT 540

AATCCACAAT TAGGGCTAGA ATCAGAAATT CCAGTTGCTC CTAGATCTAA ACGTCAAACA 600

AGACAGGCTG AAGAAGTACA GGCAAAAGAT CCTTATTTAG ATTCAGTTAA AGAACTTGAC 660

GATGTTCTTT TAAAATTTAA AAAATATTCA AAATCAATGA GTTCGATTGA AAATAAGGTT 720

TTTAGTAGTT CGGGTGGTTG TTTTAAATCA AAG 753

(2) INFORMATION FOR SEQ ID NO: 119:

(i) SEQUENCE CHARACTERISTICS:
(A) LENGTH: 747 base pairs
(B) TYPE: nucleic acid
(C) STRANDEDNESS: double
(D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 119:

TGATTTTAGC TGTTTTGTAA mCCAAAAGTG GATTATAATA ATTGGGCCTA CTAGTCTGAA 60

TCCTAGAGTC AATAAAACTT ACACTAATTG TATCTTGCGG CAATTTCGTA TTCCTCCTTT 120

AAAATTTCGA TTGCTTTTAC ACTAGCATTG AATGCTATAG ATGCACTGTA TGCATGGTTG 180

CTATATTTTG TGCCTAAATT AATCAGTCCA ACTGTTTGCA TATTAGATGT TGGGTAAATG 240

TAGAAGTTAA TTTTATTAAT ATATTCGGGT TGTAGACTGG GCAAAGTATA CTTATGAGCT 300

TTATTGTGTA GAAAGTCACT AAGCATACTA TAAAGCATTA ACATGCGTGA ATTAGCrTCA 360

AAGTCTTTGG CGTTTAACAC TATTGCAATA ATATATATTT GAAAATTTAT ACTAAATTCC 420

AAAGCATTyT CATAAAATAC ACCkGCTyTA kAATTATGAT CAAATAGATT TTCTGTACCA 480

GCAAATTTCA ATGCTATTAT ATTTGAGCTA GCAGCTGTGA TTTTTGAAAG ATATGGATGA 540

TTGTAGGTAT TTATGATATC GCACTCAAAA TTATTTTCAG TTGCATACGC CTTAAACCCT 600

TTAAATATTT TAGTTAAATG ATTTAATACC ATATCTAAAG TGAAAATCAT TCAAGTGTTA 660

CCTTATAAGT AATCTCTGAT AACATTTTGG CTGTATCAAC AAGTGGaATT GCTGCAGTGT 720

TACTACCCCT TTTAAACTTA CTTTTGA 747

(2) INFORMATION FOR SEQ ID NO: 120:

(i) SEQUENCE CHARACTERISTICS:
(A) LENGTH: 744 base pairs
(B) TYPE: nucleic acid
(C) STRANDEDNESS: double
(D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 120:

ACACTTAATT CAAAAGTACT AAGCTTTAAC CCGGAAATCT TAAGAAGATT TGAGAATTGT 60

AAATTTTAAC CTAAAAAGCA GAACCTCATA AAAGTTTGAC TTTTACCCAT AACAGTATAT 120

AATATTAATA TGTTTTTTTT CAAATTTTTT TCAAAACATA AACCTGCTAG GAAAAAAAAT 180

TATCATAAAA TCAATCCGGA TGAATTCATT CTAATTAGCG AACATCTTAT CAATTCTTAC 240

AGCATTACTC ACCAATTACT TGGGATTATC ATGGCCTCTG GAATTCCATT AACTCATATA 300

AAAAAATCAA AACATCAAAA CTCCTTACAA TTTCAAATCT GATATATTTT CTTATACGTT 360

GAACAACGGT TTGCAAATTC AAACACATTC TCTAATTTGC TCTAACAAAA TTTCTAGGTG 420

TATTGAAAGT TTAAACAAAA ACAGATTACT ATCTATTGGT GCAGACAAAA TTAATTATGT 480

AGCAAAAAAT ATTTTTGATT TTAGAATTAC TACTAAACAA CTAAAAATTA TTCATTCTTT 540

GATTGCTAGG TCAAAAGAAA CACTACATGA AATCAGATAT AACTCTCATT CACAAAACTT 600

CTTTTTAGTT AAAACACCCT GTATTTTAAA TCTGTACCAA AAGCTCAAAT ATATCAAGTC 660

ATTCGGCACC TCTAAAGCTC AATCAAAATA ATCTAAATTA TTATCGAAAC AGCTCCAATG 720

AGCTTACATC TACTATTACA AATT 744

(2) INFORMATION FOR SEQ ID NO: 121:

(i) SEQUENCE CHARACTERISTICS:
(A) LENGTH: 721 base pairs
(B) TYPE: nucleic acid
(C) STRANDEDNESS: double
(D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 121:

AGTTTGTnTA TTCCTAGTAA CAATAACATT TCAGAATAAA GTTTTGTnAC CAAAAGTTtT 60

CTTCAGCcTC AACTTGTGTT TGTAAGTTTT TTTGTTCGCT CACCTCGATT TACCTTATGT 120

TTTTAACTTT GTTTAATATG TACTTGCAAA ATAGTTTTTC TAGTAGCAAG TAACCCTCCT 180

AAAACAAAAT CAATGTATGA ATGAGCAATA TCAGTTGAAT CTTTATCCAC TTGTTCATTT 240

GGTGTAGGTA ACATATACTT GCTAGGTTTA AACTTAATAA GCTCTGGGTT TAATGGGTAA 300

ATAAGTATTT GATGTTTTAG CAAGTTTGAA GTTTCAATGT AGACATCTTC TCTATTATTA 360

ATAGCCyTGA TAGTTTGAAT CAAAACATCC TCCCATTTTT CGCAGCTACT TGCTGCACCC 420

TGTGCTGCTG CGTATGGCTT TACGAGTTTG AGCGAaGTTG TAGGgTCAAC TATTACCATC 480

ATAGGTGTAG AAAATtCGTC TCCTAGCTCT AACTTtGAAA GTCCCGCCTC AATTTTTtCA 540

AATATTTTAT CCATTTTATC TTTATCACCA CTAGCAACTT CTTCTTTTAC TTGATGTGGC 600

ATATTAAGAA GTCCATACAT ATTGGGAAGT AGACGTTTTT GATTTTTTCC ATCTTTTTGA 660

ATTGAAACAG TGCCTGTTAG TACAAAGTGA TTAATAAGTT TAATAATCTC GCTACTTGCA 720

A 721

(2) INFORMATION FOR SEQ ID NO: 122:

(i) SEQUENCE CHARACTERISTICS:
(A) LENGTH: 720 base pairs
(B) TYPE: nucleic acid
(C) STRANDEDNESS: double
(D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 122:

CAATTTCAGC TCTTAAATTT TCTATTTTAG TTCGCATACC AGTAAGTTCA ACACTAGAAT 60

ATTGCTTAAA TGCACGTATA AATCCTAATT TTAAATTAGC ACACTCTATA TCTAATTCAC 120

TTATAACTTT CCTAGCGTTA ACTTCTGATC TAAAGGTTTG CGATAAAAGG TGTTCTAAAG 180

TATCTTCACT AATTGTTACT CTAGCGTCCT GGTTAACAAT ACTTTCTCCA CTTTCCCACT 240

TTTGTCTCAT TCTCCACACA TTTACTTTAG AAACTCCTAA TTTTTTCGCT ATTTCTCTAT 300

CATTTAACGA TCCTTCTCTA AAATACACAA CATAATCATC AAAAGGCCTT TTAACTTTTT 360

TCAAAACAAT TTCTCCTAAA ATAACAAAAT TAACAAATTG TTACTCTAAG TAGTAAAGCA 420

ATTTATTAAT TGTTAACATT AACTGATAAC TTCTTGATAT TTAGCGGGGA ATATTTGTTG 480

GCCTTTATTG ATTTAGTTCG CTGCTATTTC TATAATTTTT GATTTAGAAA TAGTAGTTCA 540

TTAATTTATT GCATATTACT ATAAAATAAT TCTACTTTTT CGGAAAAATC TTTCATTTCA 600

TTCATAAGAT TTTTACTTGT GAAAAGTCTT TTATCATAAT AGTGTATACT CAAAAATAAA 660

ATATCTCTAA ATTCTTCAAT CGCATCtATT TGAAAGTCTA ATyCTAATAC TTTTCTCCTA 720

(2) INFORMATION FOR SEQ ID NO: 123:

(i) SEQUENCE CHARACTERISTICS:
(A) LENGTH: 715 base pairs
(B) TYPE: nucleic acid
(C) STRANDEDNESS: double
(D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 123:

TAATAAAAAA CTAAAGCTGT TCAACTGCAA TTGTTGCACT TGAAATTTTT TATACTAAAA 60

TAAAATACAA ATAATTATAT TAACAAATAT CGATTTTTAT AAAAAATAAG TAAAAGTAGT 120

CTAGTTTACC TGAGTATTTA AATACTTTTA ATTGAGGATG TTTTATTTTA AAAAGGAGTG 180

TAAAACTATG TCAAAAGCTG TTGACGAAGT ATATTGCTAT TCTTGTGGCA AGATTTAAAA 240

AAGATGCTGA GATTTGTATT TCTTGCGGAG TCAGAAATAA ACAAACCGAA AACTACAATA 300

AACTTATAGT ATTTTTACTA TGCTTACTTT TTGGTTATTT AGGAGTTCAC AGATTTTATG 360

TAGGTAAAAT AGGAACTGGT CTATTATACC TATTTACATT TGGATTTTTA TATGTTGGAG 420

TTTTAATCGA TCTTATTAGA ATAACAACAA ACAAGTTTAA ATGTAATTAA AAGGATTCTT 480

TAGTAAATTT TTTATTAGTC TTGTTAAAAT TATTTTTTAA TTTTTTAAGC ACATTTTGTG 540

TGAACTGCTA TTTCTATAAT CTTTGATTTA GAAATAGCAG TTCACTAGAT AATAATAAAG 600

CTAAAATTAA TATyTtAGTA TTTAATAATT CTTGAgAAAA nGTAAAATTG GTATATGTTT 660

ACTTGTTATA AAAAAATCTA TCTGGGTAGG ACTTTTAATG TTTAATAAAA TAGTG 715

(2) INFORMATION FOR SEQ ID NO: 124:

(i) SEQUENCE CHARACTERISTICS:
(A) LENGTH: 715 base pairs
(B) TYPE: nucleic acid
(C) STRANDEDNESS: double
(D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 124:

GGAGGATTTA ATATGGAATT ATTTGATGAA AATTATTATG CAAAAGCTGT GGCAAATATC 60

ATAGGAGAAG TTAAAGATCC TATTATGTAT AAATGGTTTT CGCCCGATCA AATTGAAGAT 120

GTTGATCTAC AAATGGGATA TCAAAAAACC GTAAAATGGG ACGCGTTTTT AAATGCTAAT 180

CCTACAACAA TTGCCAATGA GGTTAATACT ATCTCAACTA TTGGATTTAG TTCTGAAGTG 240

GTAAGACTTA ATTATTTGAA ATTACAGTAT AAATTCAGAC ATTTAAAGCA GACTTCTGAG 300

AAATTTTATA CTTCAGATTC ATATATTGGG GACATTAATA ATAATTTACT TCCTTTTGCT 360

CAAGCGTATA AGCTTGCAAG TAGTGAAATT ATTAAACTTA TTAATCACTT TGTATTAACC 420

GGGACTGTTT CGATTCAAAA AGATGGGAAA AATCAAAAAC GCCTGCTTCC AAATATGTAT 480

GGGCTGCTTA ATATGCCCGA GCAGATAAAA GAAGAGGTTG CTAGTGGTGA TAAAGATAAA 540

ATGGATAAAA TCTTTGAAAA GATTGAGGCT GGACTTTCAA AGTTAGAACT GGGCGACGAA 600

TTTTCCACCC CGATGGATGG TAATAGTTGA CCCAGCAACG TCACTTAAAC TAGTAAAACC 660

ATACGCnGCA GCACAGGGTG CAGCAAGTAG TTGTGAAAAA GGGAAGATGT TTTAA 715

(2) INFORMATION FOR SEQ ID NO: 125:

(i) SEQUENCE CHARACTERISTICS:
(A) LENGTH: 714 base pairs
(B) TYPE: nucleic acid
(C) STRANDEDNESS: double
(D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 125:

GTATAAAAAG CAAAAGAAAA ACATCTTCCT TCACAGAATA GTTGCCCAAA TCCAATAATA 60

ATTCATACTG GTTGAAAAAT TTCCAAGAAA ACGCAAATTA TTATTGGTGT TCATATTGTT 120

CAACATCTAG ACCAAACCGA AGTGGAGTGG CCTTTTTCTT ACTTCATTTA GTAAGTTTTC 180

AATAATTAAA CCAACAGGTA GTATTAAAAC AAAGTTTTAA TACTACCAAA GTTTTAATGG 240

CTCCTTCAAA ACAGCAGTTT TAACCGTTTC ATTCTCTCTT CTGCTAATAG TAACTGGTCA 300

TATTTAGTCA TTCCTCTCAA AACACCAATT GATGTAGCAA CAATTATCAA ATTACTAACA 360

TTAAAAAACT AAAAATATTA TAAAATATCC AAAAATAAAA ATATTCTTAT TAATTAAATA 420

ATTAATACTA ATTATTTAAT TATAGTATTA TTGCATTATA TTATAGTTAA GGAGAATATC 480

TATGAAATAC CATATAATCG TAAGTATATT TGTTTTTCTA TTTTTAAATG CTTGCAATCC 540

AGATTCTAAT ACCAATCAAA ATAATTCTAA AAAGGAATTA AAAACAGGAA GAATCCCTAA 600

TAAACAAATA AAAAATGCCC TACTTGgATG ATTTAAAAAA TTTAATAGAA ACAGCTAGTG 660

CAGGATAAAA AAATATGAAA AAAATTAGGA AGAAGAACCT TCAAACCAAT ATGG 714

(2) INFORMATION FOR SEQ ID NO: 126:

(i) SEQUENCE CHARACTERISTICS:
(A) LENGTH: 708 base pairs
(B) TYPE: nucleic acid
(C) STRANDEDNESS: double
(D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 126:

GnATTAACTG GGAAAAATTT CTCAAGTGTA ATTTTATATT CTTTTATATA ATTTTCATTT 60

AAATCGAAAG TATCGTTCTA TGCTATCCCC TTGTTCAAGT CTTCACGTTC ATGAATGGGG 120

CCTAAAAAAT TATCATCACC GCCTATTAAT TCCAATAAGT GTTTTCATTA CCAAAATTAC 180

ATTTTTTGCA TATTTTTTAG TTTAAGTTTA TTAATTCTTT TTTTACTTTC GTTTTCTCTT 240

TTTTATTTTT TATTTTTATA AATCACTTTT ATTTAAAATA TATGTATACA TATATTTTTT 300

TTACTATCTC TTTTAATAAG CCCAGACAAA TAATTACAAA TATCACACAT TAAGCAACAC 360

ACGTGCTCTT GCAATTTTGA TTTTTTAAGC CAATCTCCAA AAAATTTTTA AAAAACTTAT 420

TATATTTATT GATTGCCCyT TAAGAAACTT TTTTTATTAT GCyTTTATTA TAACTTTGTA 480

AAyCTTtCAA TAGTTTAACT TAGATAGATC GGAAAATACC TTGnCTATTA GGGCTTTATT 540

ATAACTGTTA AATTTTGnCn TTTTTTGTTA AAATTTATAT TGCCAATGAT AAAAATACGC 600

CAATTTAAAT TAAACATTTT AATAATGTCC AAATCCTCCT TATATAAGGC ACATCATAAA 660

TCAATTTTAC TGAAAAAACA AAAAGCATAT CTAAGATTTC ACCCTATT 708

(2) INFORMATION FOR SEQ ID NO: 127:

(i) SEQUENCE CHARACTERISTICS:
(A) LENGTH: 703 base pairs
(B) TYPE: nucleic acid
(C) STRANDEDNESS: double
(D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 127:

GGCCAACATG AACTATAGCC TAAAAATCTA AAAGACTAAC TTGAATTTTC TAAAAATCGT 60

AAATTTTAAC TCAAAAATCT AAACCTGCAA AATTTTAGAT TTATTACAAA GAAGTCTATC 120

ATAAACTTCG TATAATCTTG TTTCAACTCT ATCTAATCTG GCTTTAAATT CATTACCAAA 180

GCAACTAAAT CTTTAGTTTC AAATTCAAAT ACCACTCTTT TATCTAAATT ATCTATTTTA 240

AAAACCTTTA GCAATATCTC AGTTCTTTTT TTCTACCTCA TTTTTTAGTT TAAAATTTTA 300

TTTTTTATTT TTTTTATTAA CTTATTTATG ATAAAAAATT TTATTATTTA GTAAATAATT 360

ATCATATCCT TTTATTAAAG AAGAAATATA ATCTTCTCCT TTTTTTTTAT TCTTTAATGC 420

CTTAAAATCA CCAAGCAAGG TGATAAAATC TTCCTTAGCT AATGAGTAAA GACTAGCTAT 480

AATAAAATTA TTTTCATTTT CTTTTTCTTT AAAAAATTCA TCTTCTTTAT CTAGTTTCAG 540

TATTTTATTA ACTTTTTCTT TATCAAACTT AAAATATTCT AAGTAAAGTA AATATTTAAA 600

GTTTTCGGGA TCATTTTTGG CTATCAGTAA AGAAGTATTT TTTGCAAGAT TTAAATATAA 660

AGGATTACTT AAAATTTCCT TTTCTTCGGG TTGAGGCATT GGG 703

(2) INFORMATION FOR SEQ ID NO: 128:

(i) SEQUENCE CHARACTERISTICS:
(A) LENGTH: 699 base pairs
(B) TYPE: nucleic acid
(C) STRANDEDNESS: double
(D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 128:

CCGACTCAAA ACTTTACCCT TTAAATTGCT AACTTTAACT TGAAAATACT AAACTTTAAC 60

CCGAAATGAT AAAACTTTAA TTTTTGCAAT TTTATTCTCT TGTTTTTTTT AAAACGATTA 120

GAATAATCGT TGAkCAGGTT TATTGATTAT CAATAAACCT GATCTATAAT ATTATAAGCG 180

GTTTTTGCAA GTTTAATAGG AGCTATAATA TCCATGAACA AATTATTGAT ATTCATTATT 240

TTATTAGTCT TTTCATGTAA TTTAAGTAAT TCTGATCAAA ATAATCCACT AAACATGTCA 300

AATAAAGAAA AAATAAGCGA ATATCAAATA AATGAGTCGT CAAACAAATA TTCAATTTTC 360

AAACGAAATT CAAGCGTTAA AAGATACACG TTCAACCATT ATTACTAACC AAAATGATAA 420

TATTAATTCT ACTATTAACT ACCCACCTTA TATTCAAACT ATCTTAAAAA TAGAAAAACA 480

AGTTGACGGA AATATTATTA TTAATGGGAT GACTAAAGAA AGTGGCACAG AAACTAAAAA 540

GCTTTTAGAA ATTCCAAATG GGAATATTTC TCGACTTAAA GATGCAATTC AATATGGAGG 600

AAGTTTTAGG GCTAAAGATG TTAGAGAAAA TCAAACCCAA AAAGAAAACA ACAAAGACTC 660

GCATATTCAT GTCGaCAtTT TAAAGATACA TACTTTAAT 699

(2) INFORMATION FOR SEQ ID NO: 129:

(i) SEQUENCE CHARACTERISTICS:
(A) LENGTH: 695 base pairs
(B) TYPE: nucleic acid
(C) STRANDEDNESS: double
(D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 129:

GTGGCAAATA ATAATGTATT GTAAAATTTG ATTTTTTAAA ATGGTACATT ATAATATTGA 60

TAAAGAGTAT TATCAATTAA CACTTAATTT TTGCTTTTTC ATAAATTAGA ACTTATTTGA 120

ATTTTTTAAC AAGAGAATTT AAATAGGTTC TTTTATTTTA ACAAATACAA ATTGATTTTA 180

ATTCTAAATT AGAATATATT CAATTATTGA AAAGCTTATT TAAATTATTT TAATAAGCAA 240

ATTTGATTAA ACCCTAACTT TATTAAAATA ATTTATGTAA AAAGTTGTCA AAAATAGTTT 300

TTGTTATACA TATATATATG TATGTAAATA GCTAAAAAAG TTTATTGCTA TCAAAACAAT 360

CCAATCAAGT TGGGTTTAGC TAAGTTCTTA GATAAGAGAA TTTAAATAAA CCCAACTATT 420

TTTTTGTAAA ATTTTTTGTA AAAAAGCCTG ACAAAAATAG TTTTTGCTAT ATACTTATAT 480

TTTTTACTAT AAAAGGAGTA AAAAGATGGA AAATCTTTCA AACAATAATA ATCCACAAGA 540

AAATATTCAA GGAGAGCTCA AAATGATAAG TATTAATCAA CAAAGTTTTA CTGGTTGTGA 600

AATATTTGAG GAAAAATCTT CTCCCATTAA AGAAAAAAGT AAATTAAGTA AGATAGGTAA 660

GAAATTACCA GGAATAAGTA GTCAAGAATG TTTTA 695

(2) INFORMATION FOR SEQ ID NO: 130:

(i) SEQUENCE CHARACTERISTICS:
(A) LENGTH: 684 base pairs
(B) TYPE: nucleic acid

(C) STRANDEDNESS: double
(D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 130:

CAAGCAAGGG TTAACAACTA CCTTAAAGAC AAATTTAATA AAAAAGGGAA TCTAAATTTA 60

GAGGAGTGTA ATAATAATAA TAATAATAAA GAAGAAGAAG AAGAAGACAT AAGAAATAAT 120

AAAATAGAAA AATGTCAAAT AAAAAAATAT TTCAACAAAT GTAACTTTTT ATCCGAAGAA 180

GCTAAGTCCA TTTTAGAATT AAACATTAGT AAGAATAAAA CAATTGAAAT AATAAAAATA 240

ATAAAAAAAA TTGAAACCGA CTTAACAAAA AATAAAAACA AAGTTTGTTT TAAGAAAAAG 300

CAAAAAATGT TGAAAGAAAT ACTAAGCAAA ACTAAAAAGC AATTAGAAAA AAAAGGATAT 360

GACACCAAAC AACTGAAACT CAAAATCGAA AACATATATA AAAGTTATAA AACCAAGCCC 420

CATTTTATTA TTGAAAATAA AAAATACAAA GACCTAGACA AAATAAGGCT TAAACTAGAA 480

AAATCAATTG AAATTAAAAA AGAAAGTATT ACAAAAAAAT ATATACATAT AAAAGTAAAT 540

ATTTTCAACA TACTAATAGA ACAATTGAAA AAAGrmTTGG rAATAAAAAC TTTAAAGCCA 600

ATTATAAAAA ATTATCTAAA TAGCnAAAAA ACCCTAGAAT ATGATAAAGT GTTCAATACC 660

TATTATTATG nACTATTAGA AACT 684

(2) INFORMATION FOR SEQ ID NO: 131:

(i) SEQUENCE CHARACTERISTICS:
(A) LENGTH: 673 base pairs
(B) TYPE: nucleic acid
(C) STRANDEDNESS: double
(D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 131:

GTATAAATTA TCTTTTAAAT TTATTACTAA ATCTCAAATT AGATATATAC TTTGTGTTTG 60

TGCATAAACC AACTCCAGCT TTAAACAGAT CACAAGCGCT AAAGAACTTA CTAATCTACT 120

TGCAACAAAC ACCAAAACTT TGAAATAAGA AATTCAGTCA ACCTTAATAG CAAATCAAGA 180

AAAACCAAAT CTAGATATTG ATAATTGACG TACAAAAAGA AATAGACAAC ATAATTTCAA 240

ATCTAAAAGA GTTTATTAAA ACTCTACAAA CCCTACATTA AAATCTGCTT AGCATAGCTA 300

AGTATGCTCA ACAACATCTT AAAAAATTCA ATTCCAAAAA TAAAATCTAA AAATTGCACT 360

ATTATTTAAA TATCAAAACT TAATTATTAG GAGGTAATAT TAATATGAAA AAAATTTTCA 420

CATTAATATT AATTTTTAGT TTAACGATGC AAATCTTTGC TTTAAATTTC ACTTCAAAAT 480

ATAAAATTCA GAAATATGCT GAAAGAGAAA AGGAGTTCAT TCAAAACCAG AAATTAGAAA 540

AAATTTTGAA AGACCCCGAA AAGACTAAAA AGGCTCTTTT GCAATACGAA AAAGAACAAT 600

TGATAGATCT ATGGATTCCA GTAATGTTAA ATTTATTTTT ACCTTTTGGA GTGGGGcTTT 660

TGTCCAGGAG ATT 673

(2) INFORMATION FOR SEQ ID NO: 132:

(i) SEQUENCE CHARACTERISTICS:
(A) LENGTH: 660 base pairs
(B) TYPE: nucleic acid
(C) STRANDEDNESS: double
(D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 132:

CGAGTATTTT GACTCAAAAC TTTACCCTTT AAATTGCTAA CTTTAACTTG AAAATACTAA 60

ACTTTAACCC GAAATAATAA AACTTTAACT AGAATTTTTC AAAATGATAA AACTTTAACC 120

CGAAATGATA AAACTTTAAT TTTTGCAATT TTATTCTCTT GTTTTTTTTA AAACGATTAG 180

AATAATCGTT GAkCAGGTTT ATTGATTATC AATAAACCTG ATCTATAATA TTATAAGCGG 240

TTTTTGCAAG TTTAATAGGA GCTATAATAT CCATGAACAA ATTATTGATA TTCATTATTT 300

TATTAGTCTT TTCATGTAAT TTAAGTAATT CTGATCAAAA TAATCCACTA AACATGTCAA 360

ATAAAGAAAA AATAAGCGAA TATCAAATAA ATGAGTCGTC AAACAAATAT TCAATTTTCA 420

AACGAAATTC AAGCGTTAAA AGATACACGT TCAACCATTA TTACTAACCA AAATGATAAT 480

ATTAATTCTA CTATTAACTA CCCACCTTAT ATTCAAACTA TCTTAAAAAT AGAAAAACAA 540

GTTGACGGAA ATATTAATGG GATGACTAAA GAAAGTGGCA CAGAAACTAA AAAGCTTTTA 600

GAAATTCTAA ATGGGrATAT TTCTCGATTT AAAGATGaAT TCAATATGGA GGTAGTTTTA 660

(2) INFORMATION FOR SEQ ID NO: 133:

(i) SEQUENCE CHARACTERISTICS:
(A) LENGTH: 656 base pairs
(B) TYPE: nucleic acid
(C) STRANDEDNESS: double
(D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 133:

AGCACTTTAA AGAAAAAAAA GAAGAAAGAT ACCAAAATAG AGTTGCCAAC TATTTCAACA 60

AAAATTCTGA TTCAAAAATG GGTAGTGTGC AATTGGGGGA GTGTAATAAT AATAATAATA 120

ATATAAAAGA AGAAAGAAAA ATTAACGAAA TAGAAAAGTA TCAAGTAATA AAATACTTCA 180

ACAAGTGTGA CTTTTCATGT AAAGAAATTC TTCCAGTTTT ATTAACATTA AATATTGATA 240

AAGAAAACAT AATTAAAATA ATAAAAATCC TAAAAATAAC CGAAATTAAC TCAAAAAATA 300

AAAATATACG CCCTACTAAA TCTTGTATTA AAAAAAAACA AGAAAAATTA AAGGGAATTC 360

TATGTAACAC TCAAAAAGAA TTAGAAGAAA ACGGGTACAA TCCCAAACAA TTAGAAATAA 420

ATTTTCAAAA AATATACGAA AATTACAAAT ATAAACCCCA TTTTATTATT GAAAATCATA 480

AATATAGCGA TTTAAACAAC ATAAAACGTA AATTGGAAAA GTCAtTGAAA GAAAAAAAGA 540

AATTCTCAAC AAGATTATGA aATTTAAAGA TAAACGTTTT CAATATCCTA TTGAACAACT 600

AAAAAAAGAA ACAATATTGA nTTCTAAAGC CCTTATAAAA GAATTTTGAA TACCAn 656

(2) INFORMATION FOR SEQ ID NO: 134:

(i) SEQUENCE CHARACTERISTICS:
(A) LENGTH: 652 base pairs
(B) TYPE: nucleic acid
(C) STRANDEDNESS: double
(D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 134:

TGnAATTAAT GTATCAAGCG ACTCTTTGTA TCCAGTACTT TTACTGTAAA ATGTAGTCCT 60

AACAACAGGA ACTTCACCAT TTTTACCATA CACAAAAGTT GGAAATTGCC AAAAACCAAG 120

CTTTAAATTG TGATTTTTTA TAACATTTTG AATTACTTCT ACTATGACAT TGAAATCTTG 180

ATACTTATAT CCGTATCCTT TAAGACTTTT GTCAATACGT GGCAAGTTCA TTCTTAAAGT 240

ATCCATATCA TTTAAAAAGT CTATTTCTGC TTGAATATTA TTTTGTATTT CTTGATTATT 300

ATTGTTTGAA ACATTGTTCA TGTTTTCCTC CTTTATTTAG TAATAAATAA GTATATAGCA 360

AAAACTATTT TTGTCAGGCT TTTtACAAAA ATTTTTACAA AAAAGAAGTG GGACTtAACC 420

AAACTCTTTT CTTAAAGAAT CTCGTTAAGT CCCCACTATA TTATTATTTT TTGCAAATTA 480

CTAAATAAAG GTAGTCAAAC TGAAATATGT TCAAATAACT ACGCTGTTTG TAGTGTAGCC 540

CAATTTTTAA TTAAAATCAA TCAATCTTTT ACTAAGTTAT AAAAAGTATA TTAATTTAAC 600

AAAATTAATA ATTAAAATTT AATATTTTTT TAGAAAAGTA TTTAnCTTTA AA 652

(2) INFORMATION FOR SEQ ID NO: 135:

(i) SEQUENCE CHARACTERISTICS:
(A) LENGTH: 649 base pairs
(B) TYPE: nucleic acid
(C) STRANDEDNESS: double
(D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 135:

nAATTCGAGC TCGCGTACCC AGAAAGTTCA GTTAACAAAA TGTAGTCATG ACTACCTAGT 60

GTCACTTCAA TGTTGAAAAC ATAAGTTATT GTTTTGGGAT CTCTTAAGCT TATTACAGGC 120

ATACCTTTAT CTTCACTACT AATCACTGCT CTTGTTGTAG GTTCGCTTGT AAGCTCTAGC 180

TTGCCACTAT GTAACTGCGT ACCACCAATT GAAAAATAAA CTTCTCTTAA ATCATAAAAT 240

TGCATTTTTA GACCCCCTTT TAAGCACTTA AGCTGTTTTG ATAATCAAcT ATATCTTGAG 300

TAGTAATTAC TAAAGCAACA GCATTAATGC TAAAGTTATA AGTAATAkTC ACGCyAAGTT 360

CTAATTTAAG TTGyGGkGTA GGAGAAAGAG TAAGAyTTAA ATTTTTrTAC TCwATwATCA 420

GTCCTCTATC CACAAACCTT TTCAGTAAAC ATTCAATTGC TGAAGTATAT GCATTGTCTC 480

TAGCTCCACT AAGCTGCAGT GCAGATAATT TGCTATTTTG CCTATTGTTT TTGTTCCAAA 540

TTyTAATAAG CTCAATAATC GCTTCGTTTT TTATATAGTG GTATGTAAAT TGTTCGTCTA 600

TTGyACyTCC AGCTAGGTCA ACACCTTCTT TAAAGGCAGG TAAACCATC 649

(2) INFORMATION FOR SEQ ID NO: 136:

(i) SEQUENCE CHARACTERISTICS:
(A) LENGTH: 644 base pairs
(B) TYPE: nucleic acid
(C) STRANDEDNESS: double
(D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 136:

TAAGCAAAAG GnCCTAATGT TCAnGATAAT CCACAATTAG GGTTAGAATC AGAAATTCCA 60

GTTGCTCCTA GATCTAAACG TCAAGCAAGA CAGGCTGAAG AAGCACAAGC AAAAGATCCT 120

TATTTAGATT CAGTTAAAGA ACTTGACGAT GTTCTTTTAA AATTTAAAAA ATATTCAAAA 180

TCAATGAGTT CGATTGAAAA TAAGGTTTTT AGTAGTTCGG GtGGTTGTTT TAAATCAAAG 240

AATGmGCGAG TTAATGCTTA TTCTTTTACA TrTTCAAGCT TTGCAGACAA AATAGAAGAA 300

TACCTTTATG ATCCAGCAAA TAGTTTTCCA TATAAGCGTG GGGTTAAACT TGTTCCAAAA 360

GAGAAyTCTA TATATGTTGA AGTTGGTGCT GATACTGATA TGTATGGGAT ATGTGTAGAT 420

GTATGTGAGT TTAGTAGTAC TGCGTATGTA TTACCAATTA CGAATAACTT TGAAGGGTwT 480

CTTGTTACAA GAAATCCGAG TATAAAAATG GGaGAAATAT kGGaTATAAA TAACAATGGG 540

GTTATaTCAA GGCTGGTGGT GGGCCmCCAA cCGyAATTAa TGCATATGCC CTCTCTGaTT 600

CATTACAATC AATTTTGGCA CCCGAAGATG AAGATCAAGA TCAG 644

(2) INFORMATION FOR SEQ ID NO: 137:

(i) SEQUENCE CHARACTERISTICS:
(A) LENGTH: 636 base pairs
(B) TYPE: nucleic acid
(C) STRANDEDNESS: double
(D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 137:

AAAAAGTAAT GAATCATGCA TGCAGATTAC TTATTCTTAC CATAACAATA TTTGAAATCA 60

ATTTAATATT AGAAAATTAT TCTCAAAAAA CTCTACTCAA ATTTTATAAC GAAAATCTCA 120

AAAACCGAAA TCTAACTCCT AGTGTTATAT CAACAATAGA AAAATACTTA AACCAATTAG 180

AAAAAGAAAT AAACGTCATA GTCGAATTCT ATTTTAAAGA CAATCAATCC ATAATTTATT 240

ATAAACTTAA TTACACCCTA GAAAAAGTTT GCTTAAAACT ACAAGAATAC TACAAATTAT 300

TCTACAAAAA ATTAAAACAA TTTTTACAAA AAAACACTAC TACTACTTAA TTGTAAAAAA 360

TTATATCTTT GCAAATTAAG CAAATTTAGA AATATAAATT GCAAAGATAT ATATTTTTAT 420

ATGATAAATA ATAAAAATTA CTAGGAATAC TAACTTGGAA AAACTTTTGA AAAAAATAAT 480

AAAAATGAAT TACAAAAATA AGCTATCTTC TCACTTAATA ATTCTTATTT ACACACTAGG 540

CGACACTGAA CTAAATTTAA ATATTGAGTA CTATAGTAGG GGCTTTATAC ACCACGTGTT 600

TAATTCTAAC ATACATAAAT ATTGCAATAC TACTGA 636

(2) INFORMATION FOR SEQ ID NO: 138:

(i) SEQUENCE CHARACTERISTICS:
(A) LENGTH: 632 base pairs
(B) TYPE: nucleic acid
(C) STRANDEDNESS: double
(D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 138:

CCAATAACAT GACTTTAAAG GTTGCTTATT GACAATATGT TATTTTTCAA CTTTGTTGCT 60

TGAGAACTTT TTTTTTCAAT TTGCTTGTAA AGTTCCGTTA AATGAGATTC AAGCTTAGCA 120

ATATCTGTTT TTATAGAATC CTTATCATCT TGATAATCTA ATAAAAGTTG GTTTAACATG 180

TCTGATACGA TTGATTTCAT GGATAATAAT TTTTCAAACG AATTTTTAAG CTTCTCTAAA 240

TTTGAAATTT CTAGTTTATC TAAAGCATCC TTTTTAGGAT ATAAGTGATT AATCGCTGTA 300

TCAAAAGTTC TTCCAAGGTT GCAAAAAGTG CCAAATAGGA GGGTTTTTTG TTTTGATTGA 360

ATTAAAATCT TTGAAAGATT CGCTAACTTA TTGGTATTAA TAGGATTTAA AGTAGCATAA 420

GTGAATTTTC TATAGTTTTT AGATCTATCG GTATTATCTG CTACCGTTTC ATATGATGCT 480

ACCCAATAAA TTTCTTTGAA AATTGATATT CCATATTGGT TTGAAGGTTC TTCTTCTAAT 540

TTTTTTTCAT ATTTTTTTCT ATCCTCGTTA GCTGTTTCTA TTAAATTTTT TAAATCATCA 600

AGTAGCTTAT TTTTTATTTG TTTATTAGGG AT 632

(2) INFORMATION FOR SEQ ID NO: 139:

(i) SEQUENCE CHARACTERISTICS:
(A) LENGTH: 628 base pairs
(B) TYPE: nucleic acid
(C) STRANDEDNESS: double
(D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 139:

GACTTCTTTA ATAGATCAAA AAGCTCACTT CCATCACCCC CAAGAACACT ATTAACAGCG 60

GGGATCCTCT AGAGTCGACC TGCAGGCATG CAAGCTTGGC ACTGGCCGTC GTTTTACAAC 120

GTCGTGACTG GGAAAACCCT GGCGTTACCC AACTTAtCGC CTTGCAGCAC ATCCCCCTTT 180

CGCCAGCTGG CGTAATAGCG AAGAGGCCCG CACCGATCGC CCTTCCCAAC AGTTGCGCAn 240

CTGAATGGCG AATGGCGCCT GATGCGGTAT TTTCTCCTTA CGCATCTGTG CGGTATTTCA 300

CACCGCATAT GGTGCACTCT CAGTACAATC TGCTCTGATG CCGCATAGTT AAGCCAGCCC 360

CGACACCCGC CAACACCCGC TGACGCGCCC TGACGGGCTT GTCTGCTCCC GGCATCCGCT 420

TACAGACAAG CTGTGACCGT CTCCGGGAGC TGCATGTGTC AGAGGTTTTC ACCGTCATCA 480

CCGAAACGCG CGAGAcGAAA GGGGCCTCGT GgATACGCCT AwTTTTATAG GkTAATGTCA 540

TGaTAAtAAT GGtTTCTTAg AACGTcAGGg GGCAyTTTTC GGGGGAAAAG TGCGGGGGAA 600

CCCTAATTGG TTAATTTTTC CAAAATAC 628

(2) INFORMATION FOR SEQ ID NO: 140:

(i) SEQUENCE CHARACTERISTICS:
(A) LENGTH: 621 base pairs
(B) TYPE: nucleic acid
(C) STRANDEDNESS: double
(D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 140:

ATACCGCTAA ACTCATCTAT ATCGGAACAC ACTCCATATA AATCGTCTCC ACCACCAGCC 60

TCAACTTCTA GTTCGGTTGT TCCATCTCCA AAACTAAGCT TAACACCCCG TTTATACGGA 120

TACCCTTTAG CnGGtAATTC TCTATTTTGT CTTTACTGCT AGTGCAAACC CCACCAGAAT 180

TGGAAAAAAT TAGATTTtGG TCTCTAAAAT CAATAGAATT GCTAAGCAAT CCTGAGTCTT 240

GTTGGGGATT TTTCATTAAT GCTTGAATTT CTGCAACTTT CTTATCAAAT TCTTGTTTAA 300

TTTTTGTTAT ATCACTCATT TAAAAACTCC TTTAGGCAAT ACTTGTTCTT TTATGTCTTT 360

TTAGATTTTC ATAAAATTGA ATTCGTCTTT GCTTGTATGT ATTACTTATC GCTTGTACAA 420

ATTCTGTGAA ATTAATAGGT ACAAAATTAG AATCAAGCAA ACTTGCTCTT TCTTCTGATT 480

TAATAGCAAT ATTCCCCTTA ATAGAGTCAA CAGAAGAAGA ACTGCTACTC GCATTTTTTC 540

TTAATTTAAT ATTCACTTTT GCTAAAGAAA CAAGTTGCTC TAATATCTCT CCATCGATAT 600

GACTTATGTC TGATACTTTG G 621

(2) INFORMATION FOR SEQ ID NO: 141:

(i) SEQUENCE CHARACTERISTICS:
(A) LENGTH: 608 base pairs
(B) TYPE: nucleic acid
(C) STRANDEDNESS: double
(D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 141:

TCCAACTAAT AGTTATTGAT TTGTTCTTGT AGTCAGACGA TAAGCGTTGG TCCGTATGCA 60

AAATTtCTTC CATGTGAAAA ATCTGATGGT GCTCCAAGCA GTTGTTTTGG AACGGGTGTT 120

TTTTGAATAC TTGAAGAAGA CATTATCAAA AGATCATCAT TTCTAGATAG AGTGGCTGAT 180

GATATGCTAT TTGTAAGGCG TGATTTAATT TTACTAAAAA GGTTAGAAAT ATTAGTAGAA 240

TCGTTGTTAA TTAATTTGTC CGTTATTTTA GCATAAATTG TTTCTACAAA ATCTGTATTG 300

GCTGCAAGTT CTTCGGCAAT TGTAGACTTA ATTATCTGCT TAAAATAGTC TAATCCTTCT 360

CCTTTAAATA TTTTGTCTTT AGAAGCATCT AAAAAGTTTT TAAAGGTGAT AGCATTACTG 420

CTTGCAGCTC CATCATCAAG CAGTAAAAGA TCAGTATTGT TAACGGTCGT AACCTTATTT 480

AAATCTTTTA TTTGAACCGT TTCTTCTTCA TCAATTAGTA ATTTTTCTTG ATCATCAGCC 540

ATAAAACCTC CTTAGTTGTT AAAAGTTATA ATATTGTTAC CATCTGTATT ATTAATTTTG 600

AGAACTCT 608

(2) INFORMATION FOR SEQ ID NO: 142:

(i) SEQUENCE CHARACTERISTICS:
(A) LENGTH: 591 base pairs
(B) TYPE: nucleic acid
(C) STRANDEDNESS: double

(D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 142:

CTCCAATAGC GCTAAAAGAA TTAAATATAC TTGCTACTGA TGCTAGTTTA TTTATATCTT 60

GAATAATATT TGCGAATTCC TTTAATTCAT GAGGATCCAG GGGGCTTAAA ACAGTATAAG 120

TGTGTCTTCT ATATCTTATA GATCTTTCGG TATTGTCAGA TATATCTTCA GTCCCTGGCC 180

CCCAATTCAA TCCCCTAAAA GACGTCATCC CGTAATGGTC TTCAGGTTCT TTTTCCATAC 240

TTTTTACATA TTTTTCTTTA AAATTGTAGG CCGATTCTAT TTGCTTTTTT AAATCATTAG 300

ATAGCGTATT TTTTAGTTGT TGTTTTCTTA GTTCTTCCTT CTCTTCTTCT TGTTGTTTTT 360

TCTTAAGCTC TTCTTCTTGT TGTTTTTTCT TAAGCTCTTC TTCTTGTTGT TTTTTCTTAA 420

GCTCTTCTTC TTGTTGTTTT TTCTTAAGCT CTTCTTCTTG TTGTTTTTTC TTAAGCTCTT 480

CTTCTTGTTG TTTTTTCTTA AGCTCTTCTT CTTGTTGTTT TTTCTTAAGC TCTTCTTCTT 540

GTTGTTTTTT CTTAAGCTCT TCTTCTTGTT GTTTTTTCTT AAGCTCTTCT T 591

(2) INFORMATION FOR SEQ ID NO: 143:

(i) SEQUENCE CHARACTERISTICS:
(A) LENGTH: 586 base pairs
(B) TYPE: nucleic acid
(C) STRANDEDNESS: double
(D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 143:

GTGTTAGTGT CCCCATGTGA ATGGGTGCAC TAAAAAATTA AAAAAATAAA TTTAATATAG 60

GAGGATTAAT TAATGCTTAT TAATAAAATA AAACAAGATA ATAGAACTTT AAGACCGGAG 120

ATACAAAAAT GGGGTTGTTa CTTTTTGTGT CTGCATTATT ATACAAGTCT ATTTAAGCAA 180

CGTGAATTTA ATGCATATGA AATAAATACA GCGTATTATA GATTTATAGG ACTTGGTTAT 240

ATCAAGAGCA ATTGTTTTAT TATAAATCCA TGTATGATAC TTAATTATTA CGGAATTAGA 300

AGTAGCGTGA GATATGAAAC TGCAAATTAT TTGGGTGCAG CAAATGAATT TGAAATAAGT 360

GAAGTTAAAA TCGATAAGGk TAATGGATAT CACTTTATAT CAACAAAAAA TAAAGAAATA 420

TTATATGATT CACTTGaTTT AAAGCCACGT GGAAAAATAT TTAAAGTAAC TTCmAAACGT 480

wTwTTTAAAC tGrAATAGTT TaCTAAgTTT AAGGCACTTT TaGCACATTC ATAgCTgAAT 540

TTaTTAGCAG rAGrTAGGcC GTAGGATATA ACCAATTTCA TTGGTT 586

(2) INFORMATION FOR SEQ ID NO: 144:

(i) SEQUENCE CHARACTERISTICS:
(A) LENGTH: 585 base pairs
(B) TYPE: nucleic acid
(C) STRANDEDNESS: double
(D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 144:

GTGTTTTTTA GCAATTTATT TTCAACTCTT TTTATGTTTT TGATTATTTT AATCATAGTA 60

TCTTTATCAA CATTTAATGT TAATAAAAAT GGAAGAATTT CTTTACATAA GAAGTTACTT 120

TTGTTGAAGT ACTTTATTAC TTGATATTTT TCTATTTCGT TAATCTTTCT TTCTTCTTTT 180

ATATTATTTT TATTACTTAA ACACTCCACT GAATTTACAC TACTATTTTT GGAAACATTG 240

TCTTTAAAAT GkTTATTAAC TCTAGATTTA AATCTAGAGT TTTTTyGyTC TTTAAAGTAC 300

TTGTTGATTT TCTGGTAACA yTCTTTTTTA GGATACTTTA GCTTATAGTA AATTTCAGTT 360

CCACAATTTA CACCCATrTG TTGGTAGTAA TTAGTTGTrA CTTTTAATAC TTTTTCTAAT 420

TtGTAAAGAT AATTTTtGCAT tGTTCTTAGw GTAGTGGGAG CTAGACCAkT CcTTTTtAGA 480

TTTtCryTAw AGyArTAGAG TATGTTTTGT TGCGTGTATT TCTTATCTTT TTTGGTTAGG 540

TAATCTAGCG TTGAAGTAAG AGAGATTAAT TTGTGTTGGT GTTTG 585

(2) INFORMATION FOR SEQ ID NO: 145:

(i) SEQUENCE CHARACTERISTICS:
(A) LENGTH: 575 base pairs
(B) TYPE: nucleic acid
(C) STRANDEDNESS: double
(D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 145:

GGGAGATTTA ATATGGAATT ATTTGATGAA AATTATTATG CAAAAGCTGT GGCAAATATC 60

ATAGGAGAAG TTAAAGATCC TATTATGTAT AAATGGTTTT CGCCCGATCA AATTGAAGAT 120

GTTGATCTAC AAATGGGATA TCAAAAAACC GTAAAATGGG ACGCGTTTTT AAATGCTAAT 180

CCTACAACAA TTGCCAATGA GGTTAATACT ATCTCAACTA TTGGATTTAG TTCTGAAGTG 240

GTAAGACTTA ATTATTTGAA ATTACAGTAT AAATTCAGAC ATTTAAAGCA GACTTCTGAG 300

AAATTTTATA CTTCAGATTC ATATATTGGG GACATTAATA ATAATTTACT tCCTTTkGCT 360

CAAGCGTATA AGCTTGCAAG TAGTGAAATT ATTAAACTTA TTAATCACTT TGTATTAACC 420

GGGACTGTTT CGATTCAAAA AGATGGGAAA AATCAAAAAC GCCTGCTTCC AAATATGTAG 480

GGCTGCTTAA TAGCCCGAGC AGATAAAAGA AGAGGTTGCT AGTGGTGATA AAGATAAAnG 540

GGTAAAATCT TGAAAAGATG AGGCGGACTT CAAGT 575

(2) INFORMATION FOR SEQ ID NO: 146:

(i) SEQUENCE CHARACTERISTICS:
(A) LENGTH: 571 base pairs
(B) TYPE: nucleic acid
(C) STRANDEDNESS: double
(D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 146:

TTGGTTCCCA nCnTATTTCG TTGAAATTGT GATACTTATA GCCATAACCT TTAAGATTTT 60

TATCAATCCC CGGCAAGTTC ATCCTTAGGG TTTTCATATC TCTTAAAAAG TCAATTTCTG 120

CTTGATTAAT TTCTTGTGGA TTATTGTTTT TGCGGTTTTT CATTTTTTTA CTCCGTAAGT 180

TATAATTTTC TTATATATAA ATATATAGCA AAAACTATTT TTGTCAACTT TTTTTAATAA 240

AAATTTTTGT TAAAAGACTT AGGGCTTTGC TAAATTCTCT TTTAAAAGAA CTTAGTAAAG 300

CCCTAATATT TTTACGATCC AATATTCAAG TAGGAAATAA TGAAAAATTA TTTCCTACAA 360

AACTATATTT AGTTTAGTTC AACCTTAAAT TAAAATCAAT TAATATTATT ACACTGCGGT 420

CTATAAAAAT ACAAAAATAT AAAGCTTTTA TAAAATCTTA TTTTAAAAGA ACTTATAAAA 480

ACCTATTCTC TAAATTATTT ACAAAATTCT AAAATTAGAT TTTTAGTTCT TCATATTTCC 540

TTAAAAGTTT TTTAAGAAAA TCTTTTTGAT T 571

(2) INFORMATION FOR SEQ ID NO: 147:

(i) SEQUENCE CHARACTERISTICS:
(A) LENGTH: 555 base pairs
(B) TYPE: nucleic acid
(C) STRANDEDNESS: double
(D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 147:

AAGCGTATAA GCTTGCAAGT AGTGAAATTA TTAAACTTAT TAATCACTTT GTATTAACCG 60

GGACTGTTTC GATTCAAAAA GATGGGAAAA ATCAAAAACG CCTGCTTCCA AATATGTATG 120

GGCTGCTTAA TATGCCCGAG CAGATAAAAG AAGAGGTTGC TAGTGGTGAT AAAGATAAAA 180

TGGATAAAAT CTTTGAAAAG ATTGAGGCTG GACTTTCAAA GTTAGAACTG GGCGACGAAT 240

TTTCCACCCC GATGATGGTA ATAGTTGACC CAGCAACGTC ACTTAAACTA GTAAAACCAT 300

ACGCAGCAGC ACAGGGTGCA GCAAGTAGTT GTGAAAAATG GGAAGATGTT TTAATTCAAA 360

CTATTAAGGC TATTAATAAT AGAGAAGATG TTTACATTGA AACTTCAAAC TTGCTGAAAC 420

ATAAAATACT CATTTATCCA CTAAATTCTG AACTTATTAA ATTTAAACCT AGCAAGTATA 480

TGCTACCTAC ACCGAATGAA CAAGTTGATA AAGACTCAAC CGATGTAGCT CTTCTACATT 540

GATTTTGTTT TGGGC 555

(2) INFORMATION FOR SEQ ID NO: 148:

(i) SEQUENCE CHARACTERISTICS:
(A) LENGTH: 549 base pairs
(B) TYPE: nucleic acid
(C) STRANDEDNESS: double
(D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 148:

GTTTCTTCTT GGAGAATTTT GATTTGAAGA TTTTGAATTT TGAGATTCAT TTTCAAGATT 60

TTGGTTATTT TCTGATGGAT TTTTTGTTGA ATTTCCTGTT AAATTTTCTG AATTGGTGTG 120

ATTGCTTGTG TTTTTTAGAT TTCTAGAATT GTTGCTTCGT TTTGTTTTTT TTAGACTTTT 180

AGAAGTGGTA GGATTTTTTG GTTCGTTTGG GTTAACATTG CCAAAAGGTG CACATGATAT 240

GCAAATTGAA GTTAATATTG CTGTAATAAC GTTAAGTTTA ATAATATTTA ATTTAAAGTT 300

TTTCAAAATA TTCTCCTTAT AAATTTGAAT TAATATTTAT TAATTTTAGT TCAAATATAT 360

AATATTACAA TTTAATATCA ATATCAAATA AGTTTAATAT TATTGATATT GAAAATTAAT 420

TTCATGAGTT TTAGCGGGAT TAGATGCATG AATTTAAAAA TAAAAGTAAT CTCCCTTTTA 480

AAATATGAAG TGnAACAATT GTTGGATTAA GGGGTTAATC CAGGAnCAGA GGGAATTAGA 540

AATATAACG 549

(2) INFORMATION FOR SEQ ID NO: 149:

(i) SEQUENCE CHARACTERISTICS:
(A) LENGTH: 544 base pairs
(B) TYPE: nucleic acid
(C) STRANDEDNESS: double
(D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 149:

TTTTGTAATT TTCATATTCA TTAATTAAAT GATAAGACTT CTTTTTTTAA TGaAAAATAA 60

TATTTCAAAA ATAAAATAAG CTCTTTTAGT ATCTTCTTTA CAAAATTCGT AAAACCCTTT 120

GTTTTTTATT AAAATCCTAA TAGACATTTT TCTATTATTT ACTTCAGGCA AATTTTCTTT 180

TTTTTACGTT TGATAAATTC TCTTTTAACT GTCTTTGTAT TCCTCTAAAG CCCTATTTTT 240

TAATTTTATC TATATAAACA GGCCTCCTCT AAAACCCTTT TTCCGTAAAC TTTTTTTGCT 300

ATACTTTGTA TTAATAAATT TCCTAAAAGG AATTTTAAAA TTTTCTTTTA TTAAATCTTT 360

AAATACACTT TGATCTTCTT TTATACAAAG GGGAAATACA TGGCTTGATT CACTGCATCT 420

ATTTGAAACA AAACATCTCT ATAAAATTCT AGAGGTAAAA GAATGAAAAA AATTATTTGT 480

CGATAAAAAT CCTGTTGTAA TAAGGGTTAT ATTATTCTCn ATTTCCnAAA TATATTTTTT 540

TATA 544

(2) INFORMATION FOR SEQ ID NO: 150:

(i) SEQUENCE CHARACTERISTICS:
(A) LENGTH: 533 base pairs
(B) TYPE: nucleic acid
(C) STRANDEDNESS: double
(D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 150:

GAGACTTTGA AGCTATCTCG TCAGGGGTAG TGGAGTCAAT CTTGAAATAC CACCCTTGTT 60

TAATTAGGTT TCTAACTTAT AGAAATATGA GGAGAGTGCC AGGTGGGTAG TTTGACTGGG 120

GCGGTCGCCT CCTAAAGAGT AACGGAGGTG CGCAAAGGTT ACCTTAGAGT GGTTGGAAAT 180

CACTCTGTAA GTGTAAAGGC ATAAGGTAGC TTAACTGTAA GACTGACAAG TCGAACAGAT 240

ACGAAAGTAG GTCTTAGTGA TCTGGCGGTG GCAAGTGGAA GCGCCGTCAC TTAACGAATA 300

AAAGGTACTC CGGGGATAAC AGGCTTATCC TTCCCAAGAG TTCACATCGA CGGAAGGgTT 360

TGGCACcTCG ATGTCGGCTC ATCGCATCCT AGGGcTGGAG CAGgTCCTAA GGGTATGGCT 420

GTTCGCCATT TAAAGCGGTA CCGAGCTGGG TTCAGAACGT CGTGAGACAG TTTGGTCCCT 480

ATCTGCCACA AGCGTTGGAT ATTGAGAGGA GCTATnTTAG TACGAGAGGA CCG 533

(2) INFORMATION FOR SEQ ID NO: 151:

(i) SEQUENCE CHARACTERISTICS:
(A) LENGTH: 521 base pairs
(B) TYPE: nucleic acid
(C) STRANDEDNESS: double
(D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 151:

TGGTTTTGAA GCTTTTTTAG TAGGCTTAGA AGAAATTTTT AGTGAATTTT TAAGAATTTT 60

ATTTTCATTT AGCACATTTT GATAATCTTG AAATAGTTTG AGCATAAAAT CCATGTTGAA 120

ATTATTTAAA TTAAAATAAT TATTAGTGTT CATAAAATCC TCTCCTTGAA GGTGTTACTT 180

TTAAATTAAG TAAAAGTAAT AAAAATAGAT AAAAATAGTA ATTTATATTG TACCAAAAAC 240

GAAAAATTTT AGTCAAATTT TGTGAGTTCT CATTGCATGA GAAATTTGGG TTGTAGGGAG 300

GCTGTTATAA ATAGAATGGG CATTTTCTGA GGGTGTCGGC TAAGAAAGAC TACATACTTT 360

AGCTAATATA TAGCAAAGAC TTTGAAATTT AATTTGTATG TGTTTTATAG TCTTTTGTAA 420

TGAGTAGTGC ATTTGCAATG GAGAGATTTT GGGGAGTTGT TTAAAATTAC ATTTGCGTTT 480

TGTTAAAATG TAACAGCTGA ATGTAACAAA ATTATATATT T 521

(2) INFORMATION FOR SEQ ID NO: 152:

(i) SEQUENCE CHARACTERISTICS:
(A) LENGTH: 501 base pairs
(B) TYPE: nucleic acid
(C) STRANDEDNESS: double
(D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 152:

GTTCTCAAAT TTTTATAAAT CTTGTTTAGG AATTTTCTTT TTTTCTTTCA ATTGACTTTT 60

CTAATTTACG CTTTATGTAA CTTAGaTCGC TATATTTATG ATTTTCAATA ATAAAGTGGG 120

GTTTATATTT GTAATTTTCG TATATTTTTT GAAAATTTGT TTCTAATTGT TCTGAATTGT 180

ATCCACTTTT TTCTAATTCT TTTTGAGTGT TGCATAGAAT TTTCTTTAAT TTTTCTTGTT 240

TTTCTTTACT GCAAGATTTA GGAAAGTAAA TATTTTTATT TTTTGCTTTA ATATCAGTTC 300

TTTTTATGGT TTTAATTATT TTGATCATAG TATCTTaTCA ACATTTAAAT TTAATAAAAT 360

TGAAAGAATT tCTTTACATG AAAAGTTGCA TTtATtGAAA TAATTtATtA CTGATACTTT 420

CTATTcATTA AcCTTCTTCT cTTTATAtAt TTTATTACTA TTACAGGAnT CACACGTACA 480

CTACCCATTT TGAACCGAAT T 501

(2) INFORMATION FOR SEQ ID NO: 153:

(i) SEQUENCE CHARACTERISTICS:
(A) LENGTH: 488 base pairs
(B) TYPE: nucleic acid
(C) STRANDEDNESS: double
(D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 153:

GTTCCnCnTT ATCAAGGAAA TTAACGACAT CTTTCATTAT GGTTTCATTT GCAAATTTAA 60

CAAAATTGTC TTTTTCTTTT TCTAATCTTT TTCTGAAGTT TTCAAATTCT GCTTGTTTTC 120

TTAAATACAA ATCTTTAAGA TTGGAGATTT CATTTTCAAG TTCAGCAATT TTTTTATCAG 180

AATTTACTAA ATTTAAGTTT TCTTTTTTTT GAGATTTTGT ATTTTTATTA TCTTGTTTGT 240

TGTTTTTTTC AGATTCGCTT TTAGTTTCTT TTTTTTCCAT TTTTCCTCCT kTGATAAAGC 300

ATTTTATCTT TAAAAAAAAT ATTTTACAAA TTTTnTTCTT TCytGAAATT TAAnAAAATG 360

GAGTCATTTT GnGGCATTTG TAAGATGTAG ATTTTTCTTA AGCTTTCAGT AAGAGTGTTA 420

TATGnATACA TAGGTTATTT AGTnAAAATG TTCGTGTGTA TTTTGTGTCA AAAGAAAAAA 480

TTTAAGTT 488

(2) INFORMATION FOR SEQ ID NO: 154:

(i) SEQUENCE CHARACTERISTICS:
(A) LENGTH: 459 base pairs
(B) TYPE: nucleic acid
(C) STRANDEDNESS: double
(D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 154:

GGAGAGGCGG TTTGCGTATT GGGCGCTCTT CCGCTTCCTC GCTCACTGAC TCGCTGCGCT 60

CGGTCGTTCG GCTGCGGCGA GCGGTATCAG CTCACTCAAA GGCGGTAATA CGGTTATCCA 120

CAGAATCAGG ATAACGCAGA AAGAACATGT GAGCAAAAGG CCAGCAAAAG GCCAGGAACC 180

GTAAAAAGGC CGCGTTGCTG GCGTTTTTCC ATAGGCTCCG CCCCCCTGAC GAGCATCACA 240

AAAATCGACG CTCAAGTCAG AGGTGGCGAA ACCCGACAGG ACTATAAAGA TACCAGGCGT 300

TTCCCCCTGG AAGCTCCCTC GTGCGCTCTC CTGTTCCGAC CCTGCCGCTT ACCGGATACC 360

TGTCCGCCTT TCTCCCTTCG GGAAGCGTGG CGCTTTCTCA TAGCTCACGC TGTAGTATCT 420

CAGTTCGGTG TAGTCGTTCG CTCCAAGCTG GGCTGTGTG 459

(2) INFORMATION FOR SEQ ID NO: 155:

(i) SEQUENCE CHARACTERISTICS:
(A) LENGTH: 368 base pairs
(B) TYPE: nucleic acid
(C) STRANDEDNESS: double
(D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 155:

GAACAAGAGT CGAAAGTAGG TGTTAGTGAT sTGGCGGTGG CAAGTGGAAG CGCCGTCACT 60

TAACGAATAA AAGGTACTCC GGGGATAACA GGCTTATCCT TCCCAAGAGT TCACATCGAC 120

GGAAGGGTTT GGCACCTCGA TGTCGGCTCA TCGCATCCTA GGGCTGGAGC AGGTCCTAAG 180

GGTATGGCTG TTCGCCATTT AAAGCGGTAC GCrAGCTGGG TTCAGAACGT CGTGAGACAG 240

TTTGGTCCCT ATCTGCCACA AGCGTTGGAT ATTTGAGAGG AnCTATCTTT AGTACGAGAG 300

GACCGAGATG GACGAACCTC TAGTGTGCCA GTTATCCTGC CAAGGGTAAG TGCTGGGTAG 360

CTACnTTC 368

Applicant's or agent's file reference number	PB370PCT	International application No.	Unassigned

INDICATIONS RELATING TO A DEPOSITED MICROORGANISM

(PCT Rule 13*bis*)

A. The indications made below relate to the microorganism referred to in the description on page 8, line 23.

B. IDENTIFICATION OF DEPOSIT — Further deposits are identified on an additional sheet ☐

Name of depositary institution

American Type Culture Collection

Address of depositary institution (*including postal code and country*)

10801 University Boulevard
Manassas, Virginia 20110-2209
United States of America

Date of deposit	Accession Number
August 8, 1998	**202012**

C. ADDITIONAL INDICATIONS *(leave blank if not applicable)* — This information is continued on an additional sheet ☐

D. DESIGNATED STATES FOR WHICH INDICATIONS ARE MADE *(if the indications are not for all designated States)*

E. SEPARATE FURNISHING OF INDICATIONS *(leave blank if not applicable)*

The indications listed below will be submitted to the International Bureau later (*specify the general nature of the indications, e.g., "Accession Number of Deposit"*)

For receiving Office use only	For International Bureau use only
☐ This sheet was received with the international application	☐ This sheet was received by the International Bureau on:
Authorized officer	Authorized officer

What Is Claimed Is:

1. An isolated nucleic acid molecule comprising a polynucleotide having a nucleotide sequence selected from the group consisting of:

(a) any one nucleotide sequence of SEQ ID NOS:1-155; or

(b) a nucleotide sequence complementary to any one of the nucleotide sequences in (a).

(c) a nucleotide sequence at least 95% identical to any one of the nucleotide sequences of SEQ ID NOS:1-155; or,

(d) a nucleotide sequence at least 95% identical to a nucleotide sequence complementary to any one of the nucleotide sequences of SEQ ID NOS:1-155.

2. An isolated nucleic acid molecule of claim 1 comprising a polynucleotide which hybridizes under stringent hybridization conditions to a polynucleotide having a nucleotide sequence identical to a nucleotide sequence in (a) or (b) of claim 1.

3. An isolated nucleic acid molecule of claim 1 comprising a polynucleotide which encodes an epitope-bearing portion of a polypeptide in (a) of claim 1.

4. Computer readable medium having recorded thereon the nucleotide sequence depicted in SEQ ID NOS:1-155, a representative fragment thereof or a nucleotide sequence at least 95% identical to a nucleotide sequence depicted in SEQ ID NOS:1-155.

5. A method for making a recombinant vector comprising the step of inserting an isolated nucleic acid molecule of claim 1 into a vector.

6. A recombinant vector produced by the method of claim 5.

7. A host cell comprising the vector of claim 6.

8. A method of producing a polypeptide comprising:

(a) growing the host cell of claim 7 such that the protein is expressed by the cell; and

(b) recovering the expressed polypeptide.

9. An isolated polypeptide comprising a polypeptide selected from the group consisting of:

(a) a polypeptide encoded by an ORF of any one sequence of SEQ ID NOS:1-155;

(b) a polypeptide encoded by an ORF of any one sequence of SEQ ID NOS:1-155 except the N-terminal residue;

(c) a fragment of the polypeptide of (a) having biological activity; and

(d) an epitope-bearing fragment of the polypeptide of (a).

10. An isolated antibody specific for the polypeptide of claim 9.

11. A polypeptide produced according to the method of claim 8.

12. An isolated polypeptide comprising an amino acid sequence at least 95% identical to a sequence selected from the group consisting of an amino acid sequence of any one of the polypeptides in Table 1.

13. An isolated polypeptide antigen comprising an amino acid sequence of an *B. burgdorferi* epitope shown in Table 4.

14. An isolated nucleic acid molecule comprising a polynucleotide with a nucleotide sequence encoding a polypeptide of claim 9.

15. A host cell which produces an antibody of claim 10.

16. A vaccine, comprising:

(1) one or more *B. burgdorferi* polypeptides selected from the group consisting of a polypeptide of claim 9; and

(2) a pharmaceutically acceptable diluent, carrier, or excipient;

wherein said polypeptide is present, in an amount effective to elicit protective antibodies in an animal to a member of the *Borrelia* genus.

17. A method of preventing or attenuating an infection caused by a member of the *Borrelia* genus in an animal, comprising administering to said animal a polypeptide of claim 9, wherein said polypeptide is administered in an amount effective to prevent or attenuate said infection.

18. A method of detecting *Borrelia* nucleic acids in a biological sample comprising:

(a) contacting the sample with one or more nucleic acids of claim 1, under conditions such that hybridization occurs, and

(b) detecting hybridization of said nucleic acids to the one or more *Borrelia* nucleic acid sequences present in the biological sample.

19. A method of detecting *Borrelia* nucleic acids in a biological sample obtained from an animal, comprising:

(a) amplifying one or more *Borrelia* nucleic acid sequences in said sample using polymerase chain reaction, and

(b) detecting said amplified *Borrelia* nucleic acid.

20. A kit for detecting *Borrelia* antibodies in a biological sample obtained from an animal, comprising

(a) a polypeptide of claim 9 attached to a solid support; and

(b) detecting means.

21. A method of detecting *Borrelia* antibodies in a biological sample obtained from an animal, comprising

(a) contacting the sample with a polypeptide of claim 9; and

(b) detecting antibody-antigen complexes.

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ABSTRACT

Borrelia burgdorferi Polynucleotides and Sequences

The present invention provides polynucleotide sequences of the genome of *Borrelia burgdorferi*, polypeptide sequences encoded by the polynucleotide sequences, corresponding polynucleotides and polypeptides, vectors and hosts comprising the polynucleotides, and assays and other uses thereof. The present invention further provides polynucleotide and polypeptide sequence information stored on computer readable media, and computer-based systems and methods which facilitate its use.

Figure 1

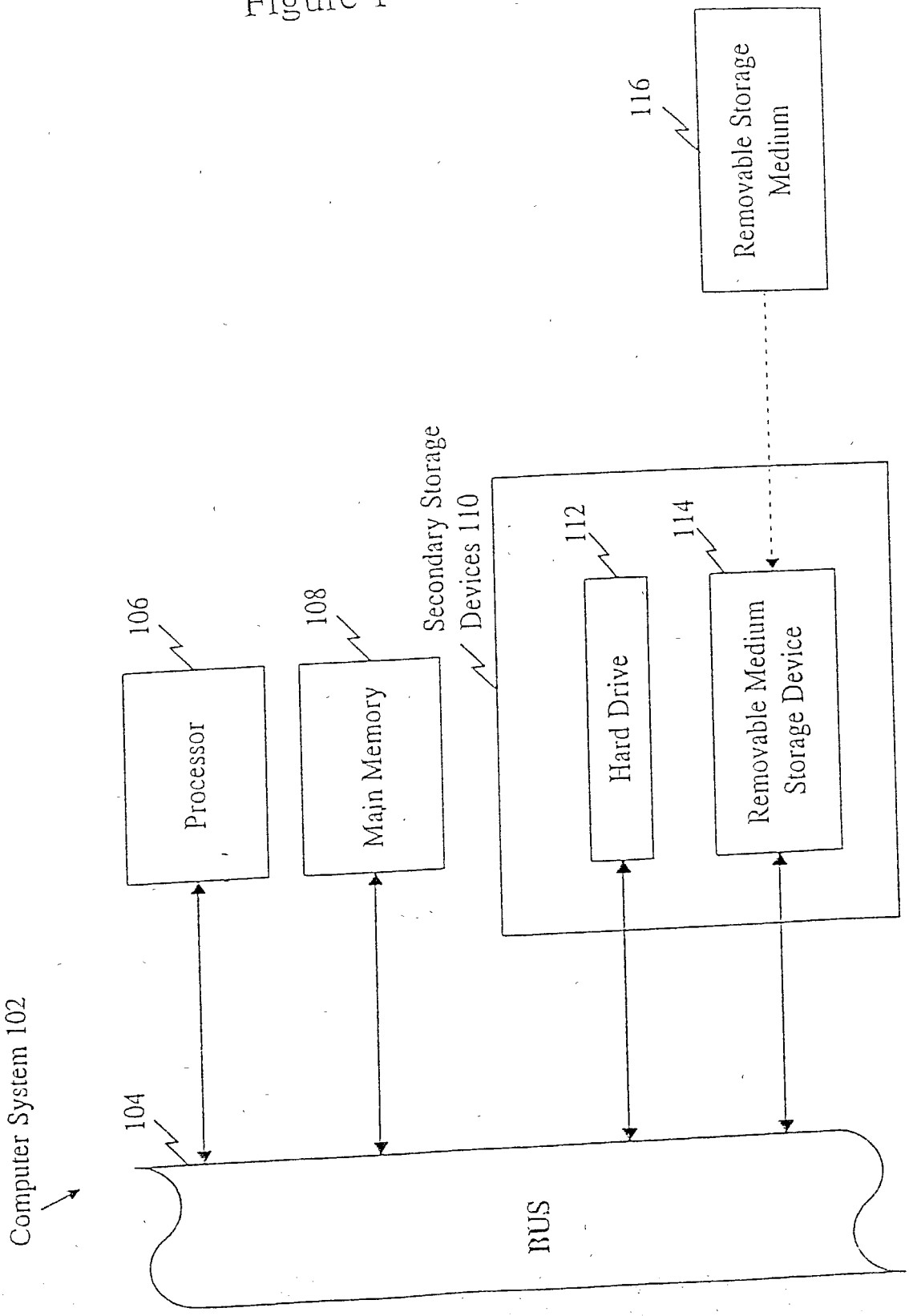

Computer System 102
104
BUS
106
Processor
108
Main Memory
Secondary Storage Devices 110
112
Hard Drive
114
Removable Medium Storage Device
116
Removable Storage Medium

Figure 2

AB 373 and 377
DNA Sample Files
(Macintosh)
Vector Removal
End Trimming
[Factura™]
Write Sequence Data &
Features to Database
[Loadis]
Treponema pallidum
Relational Database
(Unix)
ANNOTATION
Open Reading Frame
Detection
[zorf]
Nucleotide, Protein Sequence
Comparisons
[BLASTN, BLASTP]
Sequence retrieval
and ambiguity filter
[extrseq, seq_filter]
Write Data to
T. pallidum database
[lassie]
ASSEMBLY
Rapid Assembly
and Ordering of
Thousands of Sequences
[TIGR Assembler]
[asm_align]

OIPE JC176 SEP 13 2002 PATENT & TRADEMARK OFFICE

#6

PTO/SB/01 (10-01)
Approved for use through 10/31/2002 OMB 0651-0032
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Under the Paperwork Reduction Act of 1995, no persons are required to respond to a collection of information unless it contains a valid OMB control number

DECLARATION FOR UTILITY OR DESIGN PATENT APPLICATION (37 CFR 1.63)

[ ] Declaration Submitted with Initial Filing OR [x] Declaration Submitted after Initial Filing (surcharge (37 CFR 1.16 (e)) required)

Attorney Docket Number	PB370US
First Named Inventor	Claire M. Fraser
COMPLETE IF KNOWN	
Application Number	09/830,228
Filing Date	April 24, 2001
Art Unit	Not Yet Assigned
Examiner Name	Not Yet Assigned

As the below named inventor, I hereby declare that:

My residence, mailing address, and citizenship are as stated below next to my name.

I believe I am the original and first inventor of the subject matter which is claimed and for which a patent is sought on the invention entitled:

Borrelia burgdorferi Polynucleotides and Sequences

(Title of the Invention)

the specification of which

[ ] is attached hereto

OR

[x] was filed on 04/24/2001 as United States Application Number or PCT International Application No. 09/830,228 and was amended on _____ *(if applicable).*

I hereby state that I have reviewed and understand the contents of the above identified specification, including the claims, as amended by any amendment specifically referred to above.

I acknowledge the duty to disclose to the United States Patent and Trademark Office all information known to me to be material to patentability as defined in 37 C.F.R. § 1.56.

I hereby claim foreign priority benefits under 35 U.S.C. 119(a)-(d) or (f), or 365(b) of any foreign application(s) for patent, inventor's or plant breeder's rights certificate(s), or 365 (a) of any PCT international application which designated at least one country other than the United States of America, listed below and have also identified below, by checking the box, any foreign application for patent, inventor's or plant breeder's rights certificate(s), or of any PCT international application having a filing date before that of the application on which priority is claimed.

Prior Foreign Application Number(s)	Country	Foreign Filing Date (MM/DD/YYYY)	Priority Not Claimed	Certified Copy Attached? YES	NO
			[ ]	[ ]	[ ]
			[ ]	[ ]	[ ]
			[ ]	[ ]	[ ]
			[ ]	[ ]	[ ]

[ ] Additional foreign application numbers are listed on a supplemental priority data sheet PTO/SB/02B attached hereto:

OIPE JC176 SEP 13 2002 PATENT & TRADEMARK OFFICE

PTO/SB/01 (10-01)
Approved for use through 10/31/2002 OMB 0651-0032
U S Patent and Trademark Office, U S DEPARTMENT OF COMMERCE
Under the Paperwork Reduction Act of 1995, no persons are required to respond to a collection of information unless it contains a valid OMB control number.

DECLARATION — Utility or Design Patent Application

Direct all correspondence to [X] Customer Number or Bar Code Label 22195 OR [ ] Correspondence address below

I hereby appoint the Practitioners at the above-listed Customer Number as my attorney(s)/agent(s) to prosecute the application identified above, including any continuation or divisional applications, and to transact all business in the United States Patent and Trademark Office connected therewith.

Name		
Address		
City	State	ZIP
Country	Telephone	Fax

I hereby declare that all statements made herein of my own knowledge are true and that all statements made on information and belief are believed to be true; and further that these statements were made with the knowledge that willful false statements and the like so made are punishable by fine or imprisonment, or both, under 18 U.S.C. 1001 and that such willful false statements may jeopardize the validity of the application or any patent issued thereon.

NAME OF SOLE OR FIRST INVENTOR: [ ] A petition has been filed for this unsigned inventor

1-

Given Name (first and middle [if any]): Claire M.		Family Name or Surname: Fraser	
Inventor's Signature: Claire M Fraser			Date: 7/9/02
Residence: City: Potomac	State: MD	Country	Citizenship: US
Mailing Address: 11915 Glen Mill Road			
City: Potomac	State: MD	ZIP: 20854	Country

NAME OF SECOND INVENTOR: [ ] A petition has been filed for this unsigned inventor

2-

Given Name (first and middle [if any]): Owen R.		Family Name or Surname: White	
Inventor's Signature: Owen White			Date: 7/2/02
Residence: City: Rockville	State: MD	Country	Citizenship: US
Mailing Address: 17816 Milk Creek Drive			
City: Rockville	State: MD	ZIP: 20855	Country

[X] Additional inventors are being named on the 2 supplemental Additional Inventor(s) sheet(s) PTO/SB/02A attached hereto

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DECLARATION

ADDITIONAL INVENTOR(S)
Supplemental Sheet

Name of Additional Joint Inventor, if any:		☐ A petition has been filed for this unsigned inventor	
Given Name (first and middle [if any]) Sherwood		**Family Name or Surname** Casjens	
Inventor's Signature			**Date**
Residence: City Salt Lake City	**State** UT	**Country**	**Citizenship** US
Mailing Address: 1360 2nd Ave.			
City Salt Lake City	**State** UT	**ZIP** 84103	**Country**

Name of Additional Joint Inventor, if any:		☐ A petition has been filed for this unsigned inventor	
Given Name (first and middle [if any])		**Family Name or Surname**	
Inventor's Signature			**Date**
Residence: City	**State**	**Country**	**Citizenship**
Mailing Address:			
City	**State**	**ZIP**	**Country**

Name of Additional Joint Inventor, if any:		☐ A petition has been filed for this unsigned inventor	
Given Name (first and middle [if any])		**Family Name or Surname**	
Inventor's Signature			**Date**
Residence: City	**State**	**Country**	**Citizenship**
Mailing Address:			
City	**State**	**ZIP**	**Country**

Name of Additional Joint Inventor, if any:		☐ A petition has been filed for this unsigned inventor	
Given Name (first and middle [if any])		**Family Name or Surname**	
Inventor's Signature			**Date**
Residence: City	**State**	**Country**	**Citizenship**
Mailing Address:			
City	**State**	**ZIP**	**Country**

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DECLARATION FOR UTILITY OR DESIGN PATENT APPLICATION (37 CFR 1.63)

☐ Declaration Submitted with Initial Filing **OR** ☒ Declaration Submitted after Initial Filing (surcharge (37 CFR 1.16 (e)) required)

Attorney Docket Number	PB370US
First Named Inventor	Claire M. Fraser
COMPLETE IF KNOWN	
Application Number	09/830,228
Filing Date	April 24, 2001
Art Unit	Not Yet Assigned
Examiner Name	Not Yet Assigned

As the below named inventor, I hereby declare that:

My residence, mailing address, and citizenship are as stated below next to my name.

I believe I am the original and first inventor of the subject matter which is claimed and for which a patent is sought on the invention entitled:

Borrelia burgdorferi Polynucleotides and Sequences

(Title of the Invention)

the specification of which

☐ is attached hereto

OR

☒ was filed on 04/24/2001 as United States Application Number or PCT International Application No. 09/830,228 and was amended on _____ *(if applicable).*

I hereby state that I have reviewed and understand the contents of the above identified specification, including the claims, as amended by any amendment specifically referred to above.

I acknowledge the duty to disclose to the United States Patent and Trademark Office all information known to me to be material to patentability as defined in 37 C.F.R. § 1.56.

I hereby claim foreign priority benefits under 35 U.S.C. 119(a)-(d) or (f), or 365(b) of any foreign application(s) for patent, inventor's or plant breeder's rights certificate(s), or 365 (a) of any PCT international application which designated at least one country other than the United States of America, listed below and have also identified below, by checking the box, any foreign application for patent, inventor's or plant breeder's rights certificate(s), or of any PCT international application having a filing date before that of the application on which priority is claimed.

Prior Foreign Application Number(s)	Country	Foreign Filing Date (MM/DD/YYYY)	Priority Not Claimed	Certified Copy Attached? YES	NO
			☐	☐	☐
			☐	☐	☐
			☐	☐	☐
			☐	☐	☐

☐ Additional foreign application numbers are listed on a supplemental priority data sheet PTO/SB/02B attached hereto:

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DECLARATION — Utility or Design Patent Application

Direct all correspondence to [X] Customer Number or Bar Code Label 22195 OR [ ] Correspondence address below

I hereby appoint the Practitioners at the above-listed Customer Number as my attorney(s)/agent(s) to prosecute the application identified above, including any continuation or divisional applications, and to transact all business in the United States Patent and Trademark Office connected therewith.

Name		
Address		
City	**State**	**ZIP**
Country	**Telephone**	**Fax**

I hereby declare that all statements made herein of my own knowledge are true and that all statements made on information and belief are believed to be true; and further that these statements were made with the knowledge that willful false statements and the like so made are punishable by fine or imprisonment, or both, under 18 U.S.C. 1001 and that such willful false statements may jeopardize the validity of the application or any patent issued thereon.

NAME OF SOLE OR FIRST INVENTOR: [ ] A petition has been filed for this unsigned inventor

Given Name (first and middle [if any]) Claire M.		**Family Name or Surname** Fraser	
Inventor's Signature			**Date**
Potomac **Residence: City**	MD **State**	**Country**	US **Citizenship**
Mailing Address: 11915 Glen Mill Road			
Potomac **City**	MD **State**	20854 **ZIP**	**Country**

NAME OF SECOND INVENTOR: [ ] A petition has been filed for this unsigned inventor

Given Name (first and middle [if any]) Owen R.		**Family Name or Surname** White	
Inventor's Signature			**Date**
Gaithersburg **Residence: City**	MD **State**	**Country**	US **Citizenship**
Mailing Address: 886 Quince Orchard Blvd., #202			
Gaithersburg **City**	MD **State**	20878 **ZIP**	**Country**

[X] Additional inventors are being named on the 2 supplemental Additional Inventor(s) sheet(s) PTO/SB/02A attached hereto

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DECLARATION

ADDITIONAL INVENTOR(S)
Supplemental Sheet

3-00

Name of Additional Joint Inventor, if any:		☐ A petition has been filed for this unsigned inventor	
Given Name (first and middle [if any]) Rebecca		**Family Name or Surname** Clayton	
Inventor's Signature Rebecca Clayton			**Date** July 24, 2002
Residence: City Hillsboro	**State** WV	**Country**	**Citizenship** US
Mailing Address: HD-44, Box 180-A			
City Hillsboro	**State** WV	**ZIP** 24946	**Country**

4-00

Name of Additional Joint Inventor, if any:		☐ A petition has been filed for this unsigned inventor	
Given Name (first and middle [if any]) Brian A.		**Family Name or Surname** Dougherty	
Inventor's Signature Brian A Dougherty			**Date** 7/15/02
Residence: City Killingworth	**State** CT	**Country**	**Citizenship** US
Mailing Address: 10 Rosemary Lane			
City Killingworth	**State** CT	**ZIP** 06419	**Country**

Name of Additional Joint Inventor, if any:		☐ A petition has been filed for this unsigned inventor	
Given Name (first and middle [if any]) Raju		**Family Name or Surname** Lathigra	
Inventor's Signature			**Date**
Residence: City Germantown	**State** MD	**Country**	**Citizenship**
Mailing Address: 19051 Steeple Pl			
City Germantown	**State** MD	**ZIP** 20874-6165	**Country**

6-00

Name of Additional Joint Inventor, if any:		☐ A petition has been filed for this unsigned inventor	
Given Name (first and middle [if any]) Hamilton O.		**Family Name or Surname** Smith	
Inventor's Signature Hamilton O Smith			**Date** 7/11/02
Residence: City Reisterstown	**State** MD	**Country**	**Citizenship** US
Mailing Address: 13607 Hanover Pike			
City Reisterstown	**State** MD	**ZIP** 21136	**Country**

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DECLARATION

ADDITIONAL INVENTOR(S)
Supplemental Sheet
Page 1 of 2

Name of Additional Joint Inventor, if any:		☐ A petition has been filed for this unsigned inventor	
Given Name (first and middle [if any]) Rebecca		**Family Name or Surname** Clayton	
Inventor's Signature			**Date**
Takoma Park **Residence: City**	MD **State**	**Country**	US **Citizenship**
Mailing Address: 6706 B. Polar Ave			
Takoma Park **City**	MD **State**	20912 **ZIP**	**Country**

Name of Additional Joint Inventor, if any:		☐ A petition has been filed for this unsigned inventor	
Given Name (first and middle [if any]) Brian A.		**Family Name or Surname** Dougherty	
Inventor's Signature			**Date**
Killingworth **Residence: City**	CT **State**	**Country**	US **Citizenship**
Mailing Address: 10 Rosemary Lane			
Killingworth **City**	CT **State**	06419 **ZIP**	**Country**

Name of Additional Joint Inventor, if any:		☐ A petition has been filed for this unsigned inventor	
Given Name (first and middle [if any]) Raju		**Family Name or Surname** Lathigra	
Inventor's Signature			**Date** 8/29/02 (02 ~~02~~)
Germantown **Residence: City**	MD **State**	**Country**	**Citizenship** India
Mailing Address: 19051 Steeple Pl			
Germantown **City**	MD **State**	20874-6165 **ZIP**	**Country**

5-00

9/13/02 mtg Atty Reg. No. 46,789

Name of Additional Joint Inventor, if any:		☐ A petition has been filed for this unsigned inventor	
Given Name (first and middle [if any]) Hamilton O.		**Family Name or Surname** Smith	
Inventor's Signature			**Date**
Reisterstown **Residence: City**	MD **State**	**Country**	US **Citizenship**
Mailing Address: 13607 Hanover Pike			
Reisterstown **City**	MD **State**	21136 **ZIP**	**Country**

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DECLARATION

ADDITIONAL INVENTOR(S)
Supplemental Sheet
Page 2 of 2

Name of Additional Joint Inventor, if any:		☐ A petition has been filed for this unsigned inventor	
Given Name (first and middle [if any]) Sherwood		**Family Name or Surname** Casjens	
Inventor's Signature			**Date**
Salt Lake City **Residence: City**	UT **State**	**Country**	US **Citizenship**
Mailing Address: 1360 2nd Ave.			
Salt Lake City **City**	UT **State**	84103 **ZIP**	**Country**

Name of Additional Joint Inventor, if any:		☐ A petition has been filed for this unsigned inventor	
Given Name (first and middle [if any])		**Family Name or Surname**	
Inventor's Signature			**Date**
Residence: City	**State**	**Country**	**Citizenship**
Mailing Address:			
City	**State**	**ZIP**	**Country**

Name of Additional Joint Inventor, if any:		☐ A petition has been filed for this unsigned inventor	
Given Name (first and middle [if any])		**Family Name or Surname**	
Inventor's Signature			**Date**
Residence: City	**State**	**Country**	**Citizenship**
Mailing Address:			
City	**State**	**ZIP**	**Country**

Name of Additional Joint Inventor, if any:		☐ A petition has been filed for this unsigned inventor	
Given Name (first and middle [if any])		**Family Name or Surname**	
Inventor's Signature			**Date**
Residence: City	**State**	**Country**	**Citizenship**
Mailing Address:			
City	**State**	**ZIP**	**Country**

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DECLARATION FOR UTILITY OR DESIGN PATENT APPLICATION (37 CFR 1.63)

☐ Declaration Submitted with Initial Filing OR ☒ Declaration Submitted after Initial Filing (surcharge (37 CFR 1.16 (e)) required)

Attorney Docket Number	PB370US
First Named Inventor	Claire M. Fraser
COMPLETE IF KNOWN	
Application Number	09/830,228
Filing Date	April 24, 2001
Art Unit	Not Yet Assigned
Examiner Name	Not Yet Assigned

As the below named inventor, I hereby declare that:

My residence, mailing address, and citizenship are as stated below next to my name.

I believe I am the original and first inventor of the subject matter which is claimed and for which a patent is sought on the invention entitled:

Borrelia burgdorferi Polynucleotides and Sequences

(Title of the Invention)

the specification of which

☐ is attached hereto

OR

☒ was filed on 04/24/2001 as United States Application Number or PCT International Application No. 09/830,228 and was amended on _____ *(if applicable)*.

I hereby state that I have reviewed and understand the contents of the above identified specification, including the claims, as amended by any amendment specifically referred to above.

I acknowledge the duty to disclose to the United States Patent and Trademark Office all information known to me to be material to patentability as defined in 37 C.F R. § 1.56.

I hereby claim foreign priority benefits under 35 U.S.C. 119(a)-(d) or (f), or 365(b) of any foreign application(s) for patent, inventor's or plant breeder's rights certificate(s), or 365 (a) of any PCT international application which designated at least one country other than the United States of America, listed below and have also identified below, by checking the box, any foreign application for patent, inventor's or plant breeder's rights certificate(s), or of any PCT international application having a filing date before that of the application on which priority is claimed.

Prior Foreign Application Number(s)	Country	Foreign Filing Date (MM/DD/YYYY)	Priority Not Claimed	Certified Copy Attached? YES	NO
			☐	☐	☐
			☐	☐	☐
			☐	☐	☐
			☐	☐	☐

☐ Additional foreign application numbers are listed on a supplemental priority data sheet PTO/SB/02B attached hereto:

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DECLARATION — Utility or Design Patent Application

Direct all correspondence to [X] Customer Number or Bar Code Label 22195 OR [ ] Correspondence address below

I hereby appoint the Practitioners at the above-listed Customer Number as my attorney(s)/agent(s) to prosecute the application identified above, including any continuation or divisional applications, and to transact all business in the United States Patent and Trademark Office connected therewith.

Name		
Address		
City	**State**	**ZIP**
Country	**Telephone**	**Fax**

I hereby declare that all statements made herein of my own knowledge are true and that all statements made on information and belief are believed to be true; and further that these statements were made with the knowledge that willful false statements and the like so made are punishable by fine or imprisonment, or both, under 18 U S C 1001 and that such willful false statements may jeopardize the validity of the application or any patent issued thereon

NAME OF SOLE OR FIRST INVENTOR: [ ] A petition has been filed for this unsigned inventor

Given Name (first and middle [if any]) Claire M.		**Family Name or Surname** Fraser	
Inventor's Signature			**Date**
Potomac **Residence: City**	MD **State**	**Country**	US **Citizenship**
Mailing Address: 11915 Glen Mill Road			
Potomac **City**	MD **State**	20854 **ZIP**	**Country**

NAME OF SECOND INVENTOR: [ ] A petition has been filed for this unsigned inventor

Given Name (first and middle [if any]) Owen R.		**Family Name or Surname** White	
Inventor's Signature			**Date**
Gaithersburg **Residence: City**	MD **State**	**Country**	US **Citizenship**
Mailing Address: 886 Quince Orchard Blvd., #202			
Gaithersburg **City**	MD **State**	20878 **ZIP**	**Country**

[X] Additional inventors are being named on the 2 supplemental Additional Inventor(s) sheet(s) PTO/SB/02A attached hereto

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DECLARATION

ADDITIONAL INVENTOR(S)
Supplemental Sheet

Name of Additional Joint Inventor, if any:		☐ A petition has been filed for this unsigned inventor	
Given Name (first and middle [if any])	Rebecca	**Family Name or Surname**	Clayton
Inventor's Signature			**Date**
Takoma Park **Residence: City**	MD **State**	**Country**	US **Citizenship**
Mailing Address:	6706 B. Polar Ave		
Takoma Park **City**	MD **State**	20912 **ZIP**	**Country**

Name of Additional Joint Inventor, if any:		☐ A petition has been filed for this unsigned inventor	
Given Name (first and middle [if any])	Brian A.	**Family Name or Surname**	Dougherty
Inventor's Signature			**Date**
Killingworth **Residence: City**	CT **State**	**Country**	US **Citizenship**
Mailing Address:	10 Rosemary Lane		
Killingworth **City**	CT **State**	06419 **ZIP**	**Country**

Name of Additional Joint Inventor, if any:		☐ A petition has been filed for this unsigned inventor	
Given Name (first and middle [if any])	Raju	**Family Name or Surname**	Lathigra
Inventor's Signature			**Date**
Germantown **Residence: City**	MD **State**	**Country**	**Citizenship**
Mailing Address:	19051 Steeple Pl		
Germantown **City**	MD **State**	20874-6165 **ZIP**	**Country**

Name of Additional Joint Inventor, if any:		☐ A petition has been filed for this unsigned inventor	
Given Name (first and middle [if any])	Hamilton O.	**Family Name or Surname**	Smith
Inventor's Signature			**Date**
Reisterstown **Residence: City**	MD **State**	**Country**	US **Citizenship**
Mailing Address:	13607 Hanover Pike		
Reisterstown **City**	MD **State**	21136 **ZIP**	**Country**

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DECLARATION

ADDITIONAL INVENTOR(S)
Supplemental Sheet
Page 2 of 2

7-00

Name of Additional Joint Inventor, if any:		☐ A petition has been filed for this unsigned inventor	
Given Name (first and middle [if any]) Sherwood		**Family Name or Surname** Casjens	
Inventor's Signature Sherwood R Casjens			**Date** 6.21.02
Residence: City ~~Salt Lake City~~	**State** UT	**Country**	**Citizenship** US
Mailing Address: 1360 2nd Ave.			
City Salt Lake City	**State** UT	**ZIP** 84103	**Country**

Name of Additional Joint Inventor, if any:		☐ A petition has been filed for this unsigned inventor	
Given Name (first and middle [if any])		**Family Name or Surname**	
Inventor's Signature			**Date**
Residence: City	**State**	**Country**	**Citizenship**
Mailing Address:			
City	**State**	**ZIP**	**Country**

Name of Additional Joint Inventor, if any:		☐ A petition has been filed for this unsigned inventor	
Given Name (first and middle [if any])		**Family Name or Surname**	
Inventor's Signature			**Date**
Residence: City	**State**	**Country**	**Citizenship**
Mailing Address:			
City	**State**	**ZIP**	**Country**

Name of Additional Joint Inventor, if any:		☐ A petition has been filed for this unsigned inventor	
Given Name (first and middle [if any])		**Family Name or Surname**	
Inventor's Signature			**Date**
Residence: City	**State**	**Country**	**Citizenship**
Mailing Address:			
City	**State**	**ZIP**	**Country**